

STIC Search Report **ĒIC 1700**

STIC Database Tracking Number: 178833

TO: Ardith Hertzog **Location: REM 9A20**

Art Unit: 1754 **February 8, 2006**

Case Serial Number:

PCT/US04/05645 10/786,671

From: Les Henderson Location: EIC 1700 **REM 4B28 / 4A30** Phone: 571-272-2538

Leslie.henderson@uspto.gov

Search Notes					
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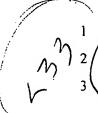


The composition of claim 6, wherein the polyoxometalate has the formula A[VkMomWnNboTaoMaXrOs] y, wherein A includes at least one counterion selected 2 3 from alkali metal cations, alkaline earth metal cations, ammonium cations, 4 quaternary ammonium cations, d-block cations, f-block cations, and combinations 5 thereof, wherein M includes at least one element selected from an f-block element 6 and a d-block element having at least one d-electron, except for vanadium, molybdenum, tungsten, niobium, or tantalum, wherein X includes at least on 7 element selected from a p-block element, a d-block element, and an f-block 8 9 element, except for oxygen, wherein k can range from 0 to 30, wherein m can 10 range from 0 to 160, wherein n can range from 0 to 160, wherein o can range from 0 to 30, where p can range from 0 to 10, wherein q can range from 0 to 30, 11 12 wherein r can range from 0 to 30, wherein s is a number so that y is greater than zero, wherein the sum of k, m, n, o, and p is greater than or equal to four; and 13 14 wherein the sum of k, m, and q is greater than zero.

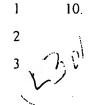


1

The composition of claim-6; wherein the polyoxometalate has the formula $[X^gV_b^{j+}M_c^{h+}Z_{12\cdot b\cdot c}^{i+}O_x]^{u-}[A]$, wherein X is at least one p-, d-, or f-block element; g is greater than or equal to 2; M is at least one f-block element or d-block element having at least one d-electron, wherein M is not vanadium; h is from 1 to 7; i is from 5 to 6; j is from 4 to 5; x is 39 or 40; Z is tungsten, molybdenum, niobium, or a combination thereof; b is from 0 to 6; c is from 0 to 6; u is from 3 to 9; and A is a counterion.



The composition of claim 6, wherein the polyoxometalate has the formula $[X^gV_b^{j+}Z_{12-b}^{i+}O_{40}]^{u-}[A]$, wherein X is at least one of phosphorus, silicon, aluminum, boron, zinc, cobalt, or iron; b is from 1 to 6, and a is from 3 to 9.



8.

9.

The Composition of claim-6, wherein the polyoxometalate has the formula $[X^{g+}M_c^{h+}Z_{12-c}^{i+}O_{40}]^{u-}[A]$, wherein X is at least one of phosphorus, silicon, aluminum, boron, zinc, cobalt, or iron; c is from 1 to 6, and a is from 3 to 9.



United States Patent and Trademark Office

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

CONFIRMATION NO. 3022

3ib Data Sheet							4110N NO. 3022	
SERIAL NUMBER 10/786,671	FILING DATE 02/25/2004 RULE	CLASS GROU 588		UP ART U 1754	JNIT	T ATTORNEY DOCKI NO. 50508-1190		
\PPLICANTS								
Nelya Okun, Alp	oharetta, GA;				\ A			
Craig L. Hill, Atl	anta, GA;		: A	H	MA			
This appln clain	. ************************************	25/2003 (PC)	ed 6	ngse	/)			
	** SMALL ENTITY **							
oreign Priority claimed .5 USC 119 (a-d) conditions n /erified and Acknowledged		STATE OR		EETS WING 0	CL	OTAL AIMS 58	INDEPENDENT CLAIMS 2	
ADDRESS 24504 FHOMAS, KAYDEN, H 100 GALLERIA PARKI STE 1750 ATLANTA, GA 30339-5948	IORSTEMEYER & RISLEY, WAY, NW	LLP						
FITLE	Is incorporating the composi	tions, and methods of usir	ng the cor	mpositions	s and m	aterials		
				☐ All Fe	es			
					1.16 Fees (Filing)			
FILING FEE FEES: Authority has been given in Paper No to charge/credit DEPOSIT ACCOUNT					1.17 Fees (Processing Ext. of time)			
RECEIVED No	for following:					1.18 Fees (Issue)		
792					Other			
		☐ Credit						

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                10141-05-6/BI OR 10421-48-4/BI OR 107-92-6/BI OR
                109-52-4/BI OR 110-81-6/BI OR 110-86-1/BI OR 13093-17-9
                /BI OR 13138-45-9/BI OR 134360-58-0/BI OR 13770-18-8/BI
                 OR 3251-23-8/BI OR 34946-82-2/BI OR 352-93-2/BI OR
                38465-60-0/BI OR 50-00-0/BI OR 503-74-2/BI OR 505-60-2/
                BI OR 57-12-5/BI OR 59858-44-5/BI OR 624-92-0/BI OR
                630-08-0/BI OR 693-07-2/BI OR 74-93-1/BI OR 7439-89-6/B
                I OR 7440-22-4/BI OR 7440-33-7/BI OR 7440-38-2/BI OR
                7440-45-1/BI OR 75-07-0/BI OR 75-18-3/BI OR 75-44-5/BI
                OR 75-50-3/BI OR 7664-41-7/BI OR 7704-34-9/BI OR
                7727-37-9/BI OR 7783-06-4/BI OR 79-09-4/BI OR 795308-36
                -0/BI OR 796042-78-9/BI)
1.3
             13 SEA ABB=ON PLU=ON L2 AND 2/NC
                D SCAN
              4 SEA ABB=ON PLU=ON L3 AND BUTANAMINIUM
                D SCAN
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L4 1.5 9 SEA ABB=ON PLU=ON L3 NOT L4

D SCAN L6

28 SEA ABB=ON PLU=ON L2 NOT L3 D SCAN

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L7 12 SEA ABB=ON PLU=ON L4 D SCAN D SCAN TI

1977 SEA ABB=ON PLU=ON POLYOXOMETAL? OR POLY(A)OXOMETAL? L8 OR POLYOXO(A) METAL? OR POLY(2A) OXO(2A) METAL?

35 SEA ABB=ON PLU=ON OKUN N?/AU L9 L10 2316 SEA ABB=ON PLU=ON HILL C?/AU L11 10 SEA ABB=ON PLU=ON L9 AND L10 2 SEA ABB=ON PLU=ON L11 AND L7 8 SEA ABB=ON PLU=ON L11 AND L8 L12 L13 D SCAN

D L13 1-8 TI AU CC

FILE 'REGISTRY' ENTERED AT 14:17:08 ON 08 FEB 2006 D SCAN L4

D L4 1-4 CN

L14 4330809 SEA ABB=ON PLU=ON (T1 OR T2 OR T3 OR LNTH OR ACTN OR SHEL OR A3 OR A4 OR A5 OR A6 OR A7) / PG AND OXO

4 SEA ABB=ON PLU=ON L4 AND L14 L15

49866 SEA ABB=ON PLU=ON L14 AND (((W OR MO OR NB)(L)O)/ELS) 1.16

L17 3 SEA ABB=ON PLU=ON L4 AND L16 1 SEA ABB=ON PLU=ON L4 NOT L17 L18 D SCAN 2458 SEA ABB=ON PLU=ON (V(L)O)/ELS AND L16 2318 SEA ABB=ON PLU=ON L19 AND 1-6/V L19 L20

D QUE L14 L21 8617 SEA ABB=ON PLU=ON L16 AND 39-40/O

1460 SEA ABB=ON PLU=ON L21 AND L19 L22 L23 2318 SEA ABB=ON PLU=ON L16 AND 6>=V

L24 47408 SEA ABB=ON PLU=ON L16 NOT 1-100/V L25 7157 SEA ABB=ON PLU=ON L24 AND 39-40/0

1652 SEA ABB=ON PLU=ON L19 AND (P OR SI OR AL OR B OR ZN L26

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OR CO OR FE)/ELS
             1112 SEA ABB=ON PLU=ON L26 AND 40/O
1435 SEA ABB=ON PLU=ON L21 AND L23
6013 SEA ABB=ON PLU=ON L24 AND 40/O
L27
L28
L29
             5008 SEA ABB=ON PLU=ON L29 AND (P OR SI OR AL OR B OR ZN
L30
                   OR CO OR FE)/ELS
             1579 SEA ABB=ON PLU=ON L20 AND L26
T.31
                   D QUE
                   D QUE L14
             1652 SEA ABB=ON PLU=ON ((P OR SI OR AL OR B OR ZN OR CO
1.32
                  OR FE) (L) V(L) (W OR MO OR NB) (L) O) /ELS AND OXO
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L33
L34
                1 SEA ABB=ON PLU=ON L4 AND L33
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1.36
             7301 SEA ABB=ON PLU=ON L30
             1317 SEA ABB=ON PLU=ON L28
L37
                  D QUE L36
             8151 SEA ABB=ON PLU=ON L35 OR L36 OR L37
L38
                3 SEA ABB=ON PLU=ON L11 AND L38
1,39
                   D SCAN
L40
              662 SEA ABB=ON PLU=ON L38 AND L8
L41
              483 SEA ABB=ON PLU=ON L38 AND (?NITRATE OR NITRIC(A)ACID)
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L42
                 6 SEA ABB=ON PLU=ON L3 AND (NITRATE OR NITRIC(A)ACID)
                3 SEA ABB=ON PLU=ON L5 NOT L42
L43
                   D SCAN
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L44
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L45
                3 SEA ABB=ON PLU=ON L44 AND L38
                  D SCAN
          429744 SEA ABB=ON PLU=ON AIR POLLUTION/SC,SX
L46
              87 SEA ABB=ON PLU=ON L38 AND L46
L47
L48
          526458 SEA ABB=ON PLU=ON TOX?/SC,SX
         20 SEA ABB=ON PLU=ON L48 AND L38

1252802 SEA ABB=ON PLU=ON PHARMACOL?/SC,SX

107 SEA ABB=ON PLU=ON L50 AND L38

QUE ABB=ON PLU=ON CONTAMIN? OR POLLUT? OR TOX? OR
T.49
L50
L51
1.52
                  POISON?
1.53
              148 SEA ABB=ON PLU=ON L38 AND L52
                  QUE ABB=ON PLU=ON PURE OR PURIF? OR CLEAN? OR
L54
                   DECONTAM?
              332 SEA ABB=ON PLU=ON L38 AND L54
L55
                 OUE ABB=ON PLU=ON WAR? OR EXPLO?
L56
L57
               77 SEA ABB=ON PLU=ON L56 AND L38
           16041 SEA ABB=ON PLU=ON L42
2027 SEA ABB=ON PLU=ON L43
181 SEA ABB=ON PLU=ON L58 AND L59
3 SEA ABB=ON PLU=ON L60 AND L38
1.58
L59
L60
L61
               9 SEA ABB=ON PLU=ON L35 AND L46
L62
L63
              10 SEA ABB=ON PLU=ON L35 AND L48
              25 SEA ABB=ON PLU=ON L35 AND L50
L64
              24 SEA ABB=ON PLU=ON L35 AND L52
44 SEA ABB=ON PLU=ON L35 AND L54
15 SEA ABB=ON PLU=ON L35 AND L57
L65
L66
L67
              67 SEA ABB=ON PLU=ON L61 OR L45 OR (L62 OR L63 OR L64
L68
                  OR L65) OR L67
              29 SEA ABB=ON PLU=ON L61 OR L62 OR L63 OR L67
9 SEA ABB=ON PLU=ON L37 AND L46
10 SEA ABB=ON PLU=ON L37 AND L48
22 SEA ABB=ON PLU=ON L37 AND L50
L69
L70
L71
L72
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24 SEA ABB=ON PLU=ON L37 AND L52
38 SEA ABB=ON PLU=ON L37 AND L54
L73
L74
              12 SEA ABB=ON PLU=ON L37 AND L57
L75
L76
              25 SEA ABB=ON PLU=ON L70 OR L71 OR L75
L77
              29 SEA ABB=ON PLU=ON L76 OR L69
              80 SEA ABB=ON PLU=ON L36 AND L46
L78
             13 SEA ABB=ON PLU=ON
105 SEA ABB=ON PLU=ON
L79
                                         L36 AND L48
L80
                                         L36 AND L50
              135 SEA ABB=ON PLU=ON
                                         L36 AND L52
L81
L82
              309 SEA ABB=ON PLU=ON L36 AND L54
L83
              66 SEA ABB=ON PLU=ON L36 AND L57
L84
               39 SEA ABB=ON PLU=ON L79 OR L77
              QUE ABB=ON PLU=ON
831 SEA ABB=ON PLU=ON
L85
                                         COMPOSIT? OR COMPN# OR COMPSN#
L86
                                         L38 AND L85
               6 SEA ABB=ON PLU=ON
                                         L86 AND L46
1.87
               68 SEA ABB=ON PLU=ON L86 AND (L48 OR L50 OR L52 OR L54
L88
                  OR L57)
L89
                3 SEA ABB=ON PLU=ON L88 AND L48
L90
               10 SEA ABB=ON
                               PLU=ON
                                         L88 AND L50
              10 SEA ABB=ON PLU=ON
1.91
                                         L88 AND L52
              42 SEA ABB=ON PLU=ON L88 AND L54
L92
L93
              11 SEA ABB=ON PLU=ON L88 AND L57
              28 SEA ABB=ON PLU=ON
1,94
                                         (L89 OR L90 OR L91) OR L93
              33 SEA ABB=ON PLU=ON L94 OR L87
66 SEA ABB=ON PLU=ON L95 OR L84
L95
L96
                  D SCAN TI
L97
               33 SEA ABB=ON PLU=ON L96 AND L85
                  QUE ABB=ON PLU=ON MIX? OR MIXT# OR MIXTURE? OR
1.98
                  BLEND? OR ADMIX? OR COMMIX?
L99
                  QUE ABB=ON PLU=ON IMMIX? OR INTERMIX? OR DOPE# OR
                  DOPING# OR DOPANT# OR IMPREGNAT? OR COMPOSIT? OR
                  COMPN#
L100
                  QUE ABB=ON PLU=ON COMPSN# OR FORMULAT? OR COMBINAT?
                  OR INTERSPER? OR AMALGAM?
              42 SEA ABB=ON PLU=ON L96 AND ((L98 OR L99 OR L100))
42 SEA ABB=ON PLU=ON L97 OR L101
QUE ABB=ON PLU=ON (MIXT# OR MIXTURE? OR BLEND? OR
T-101
L102
L103
                  ADMIX? OR COMMIX? OR IMMIX? OR INTERMIX? OR COMPOSIT?
                  OR COMPN# OR COMPSN# OR FORMULAT? OR INTERSPER?)/TI
              13 SEA ABB=ON PLU=ON L96 AND L103
42 SEA ABB=ON PLU=ON L102 OR L104
2 SEA ABB=ON PLU=ON L11 AND L105
L104
L105
L106
                  D SCAN
              8 SEA ABB=ON PLU=ON L106 OR L12 OR L13 48 SEA ABB=ON PLU=ON L107 OR L105
L107
T-108
             110 SEA ABB=ON PLU=ON L40 AND (L46 OR L48 OR L50 OR L52
T-109
                  OR L54 OR L57)
               18 SEA ABB=ON PLU=ON L109 AND L85
T.110
              32 SEA ABB=ON PLU=ON L109 AND ((L98 OR L99 OR L100))
L111
L112
               6 SEA ABB=ON PLU=ON L109 AND L103
L113
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10 SEA ABB=ON PLU=ON L114 NOT L113
L114
L115
                  D SCAN TI
L116
                8 S L11 AND L113
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L114 ANSWER 1 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:1272658 HCAPLUS

DOCUMENT NUMBER: 144:31700

TITLE: Anti penicillin-resistant streptococcus

pneumonia (prsp) heteropolyacid compounds and

their preparation method

INVENTOR(S): Liu, Shuxia; Bai, Yunpeng; Zhai, Hongju; Du,

Hongming; Li, Dehui; Liang, Dadong

PATENT ASSIGNEE(S): Northeast Normal University, Peop. Rep. China SOURCE: Faming Zhuanli Shenqing Gongkai Shuomingshu,

13 pp.

CODEN: CNXXEV

DOCUMENT TYPE: LANGUAGE:

Patent Chinese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 1616466	A	20050518	CN 2004-10011104	
				2004
				0917
PRIORITY APPLN. INFO.:			CN 2004-10011104	
				2004
				0917

The invention relates to anti penicillin-resistant streptococcus pneumonia (PRSP) polyacid compds. and their preparation method, which falls into the field of chemical synthetic drugs and their preparation method. The polyacid compound in this invention is synthesized by self-assembly of clinic drugs such as amantadine, moroxydine, 5-Fu(5-Fc) and isoniazide with polyoxometalate, where the polyoxometalate includes heteropolyacid salts of Keggin type [XM12040]n- (X = P, As, Si, Ge and M = Mo or W), Dowson type [X2M18062]n- (X = P, As, Si, Ge etc. and M = Mo, W etc.), and [LnW10036]n- (Ln = lanthanide). One or combination of more compds. prepared in this invention has outstanding curative effect on intractable streptococcus pneumonia infection which cannot be cured by antibiotics.

IT 864828-17-1P

RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(preparation and antibacterial activity of anti penicillin-resistant streptococcus pneumonia (prsp) heteropolyacid compds.)

RN 864828-17-1 HCAPLUS

CN Tungstate(4-), $[\mu 12-[orthosilicato(4-)-$

κ0:κ0:κ0:κ0':κ0':κ0':κ0'

':κ0'':κ0'':κ0''':κ0''':κ0''']]tetra

cosa-μ-oxododecaoxododeca-, tetrahydrogen, compd. with

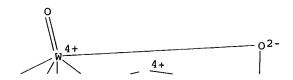
5-fluoro-2,4(1H,3H)-pyrimidinedione (1:4) (9CI) (CA INDEX NAME)

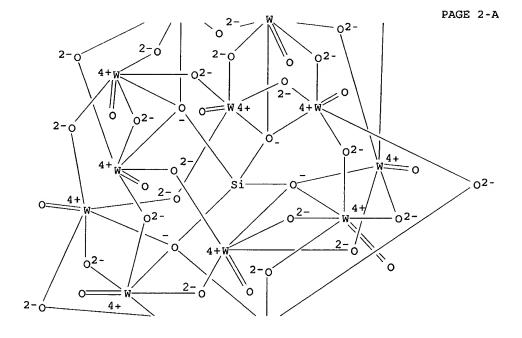
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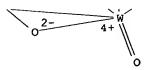
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CMF H . 1/4 040 Si W12

CCI CCS







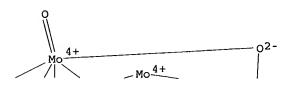
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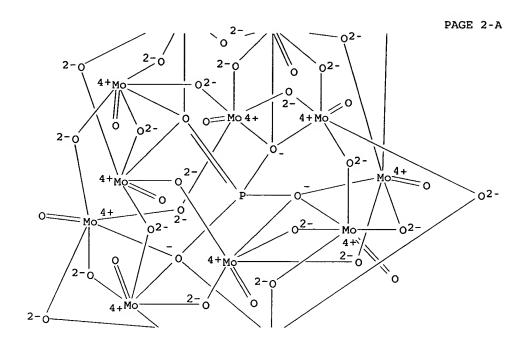
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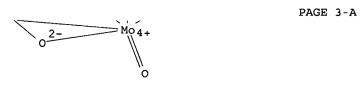
12026-57-2, H3PMo12040 12027-38-2
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation and antibacterial activity of anti penicillin-resistant streptococcus pneumonia (prsp) heteropolyacid compds.)
12026-57-2 HCAPLUS

RN

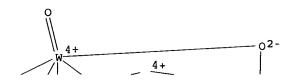
Molybdate(3-), tetracosa- μ -oxododecaoxo[μ 12-[phosphato(3-)κΟ:κΟ:κΟ':κΟ':κΟ':κΟ' ':κΟ'':κΟ'':κΟ'':κΟ''':κΟ''']]dodec a-, trihydrogen (9CI) (CA INDEX NAME)

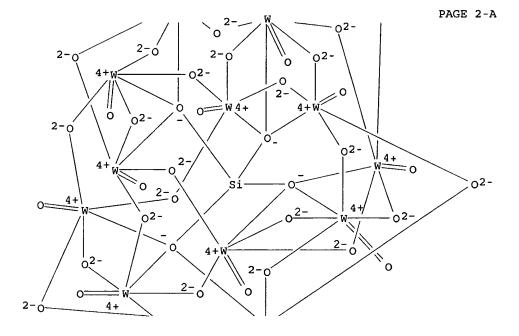






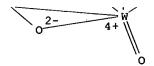
●3 H+





571-272-2538

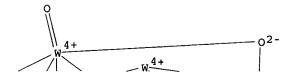
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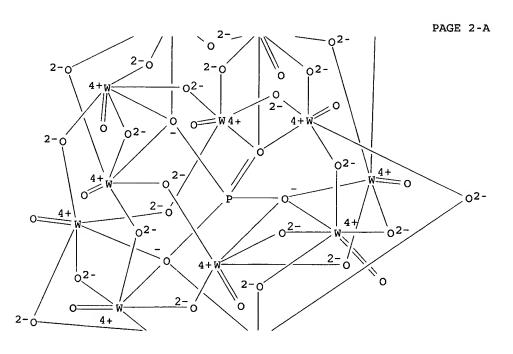


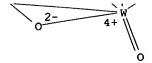
●4 H+

IC ICM C07F003-00 ICS C07F011-00; A61K031-66; A61K031-555; A61K031-28; A61P031-04 78-7 (Inorganic Chemicals and Reactions) CC Section cross-reference(s): 1, 10 ST anti penicillin resistant streptococcus pneumonia polyoxometalate prepn antibacterial activity 864828-17-1P 870470-72-7P 870470-75-0P 870470-77-2P 870470-82-9P 870470-84-1P 870470-80-7P 870470-86-3P 870470-88-5P 870470-91-0P 870470-93-2P RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation) (preparation and antibacterial activity of anti penicillin-resistant streptococcus pneumonia (prsp) heteropolyacid compds.) 51-21-8, Fluril 54-85-3, Isoniazide 768-94-5, Amantadine 3731-59-7, Moroxydine 12026-57-2, H3PMo12040 12411-74-4 12027-38-2 63055-84-5 86045-29-6 RL: RCT (Reactant); RACT (Reactant or reagent) (preparation and antibacterial activity of anti penicillin-resistant streptococcus pneumonia (prsp) heteropolyacid compds.) L114 ANSWER 2 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2005:537222 HCAPLUS DOCUMENT NUMBER: 143:371895 TITLE: Study on the photodegradation of formaldehyde with PWn/TiO2 composite catalysts AUTHOR(S): Gai, Tiejun; Lu, Xiaomeng; Deng, Qian; Xiao, Hanxi; Peng, Zhenshan Dependent of Chemistry and Chemical Engineer, CORPORATE SOURCE: Hunan University of Science & Technology, Xiangtan, 411201, Peop. Rep. China SOURCE: Huanjing Kexue Xuebao (2005), 25(5), 618-622 CODEN: HKXUDL; ISSN: 0253-2468 PUBLISHER: Kexue Chubanshe DOCUMENT TYPE: Journal LANGUAGE: Chinese PWn/TiO2 composite catalysts were prepared by the method of sol-gel via calcination technique. Structure characterization was obtained by TG-DTA, FIR, BET sp. surface area, TPR, SEM, FL. Photocatalytic performance of composite catalysts was

investigated with formaldehyde as test compound in a static photoreactor made of quartz glass. The results showed that PW12/TiO2 composite catalyst (calcinated at 350°C) kept complete Keggin structure, and the carriers effectively transferred from TiO2 to PW12, so PW12/TiO2 composite catalysts had better photocatalysis activity than pure TiO2 and that .tplbond. Ti-OH may combine with PW11 at its lacunary position, leading to structure change, so PW11/TiO2 composite catalysts had lower photocatalysis activity than PW12/TiO2. Photocatalysis reactions over two type composite catalysts accorded to L-H mechanism, which had first order kinetic equation. First-order reaction rate constant of PW12/TiO2 and PW11/TiO2 were 0.01243 min-1 and 0.005214 min-1.







●3 H+

59-6 (Air Pollution and Industrial Hygiene) Section cross-reference(s): 67 ST photodegrdn formaldehyde PW12 titania composite catalyst air purifn Air purification TΤ (photocatalytic oxidation; photodegrdn. of formaldehyde with PW12/TiO2 composite catalysts) IT 13463-67-7, Titania, uses RL: CAT (Catalyst use); USES (Uses) (photodegrdn. of formaldehyde with PW12/TiO2 composite catalysts) 1343-93-7, 12-Tungsto Phospho ric acid 12412-84-9, 11-Phosphotungstic acid RL: CAT (Catalyst use); RGT (Reagent); RACT (Reactant or reagent); USES (Uses) (photodegrdn. of formaldehyde with PW12/TiO2 composite catalysts) IT 50-00-0, Formaldehyde, processes RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); POL (Pollutant); REM (Removal or disposal); OCCU (Occurrence); PROC (Process)
(photodegrdn. of formaldehyde with PW12/TiO2 composite catalysts) TТ 5593-70-4, Tetra butoxy titanium RL: RGT (Reagent); RACT (Reactant or reagent) (photodegrdn. of formaldehyde with PW12/TiO2 composite catalysts) L114 ANSWER 3 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2005:515250 HCAPLUS TITLE: Vanadium-substituted Dawson-type polyoxometalates as versatile electrocatalysts AUTHOR (S): Keita, Bineta; Mbomekalle, Israel-Martyr; Nadjo, Louis; de Oliveira, Pedro; Ranjbari, Alireza; Contant, Roland Electrochimie et Photoelectrochimie, UMR 8000, CORPORATE SOURCE: CNRS, Laboratoire de Chimie Physique, universite Paris-Sud, Orsay, 91405, Fr. SOURCE: Comptes Rendus Chimie (2005), 8(6-7), 1057-1066 CODEN: CRCOCR; ISSN: 1631-0748 PUBLISHER: Editions Scientifiques et Medicales Elsevier DOCUMENT TYPE: Journal LANGUAGE: English A selected series of mono- and multi- V-substituted derivs. of Dawson type structure were synthesized and characterized with the aim of exploring their electrochem. and their electrocatalytic abilities. The focus was placed on the electrochem. of [P2V2W16O62]8- as a representative example. The

```
redox processes of its two V-centers were observed in a potential
     domain well pos. of those of the W-centers. They are
     pH-dependent, the first redox process exhibiting only modest and
     progressively smaller potential shifts from pH 0 to 4, while the
     second wave was far more sensitive to acidity changes from pH 0 to
     8. In contrast, mono-substituted derivs. display very small or no
     pH-dependence of the V-wave. Finally, combination of
     this diversity in the number of V atoms with the presence of As or P as the central heteroatom in these tungstic and molybdo-tungstic
     structures modulates substantially the apparent formal potentials
     that span the range from + 569 mV to + 122 mV vs SCE at pH 7.
     This leaves considerable flexibility in the choice of POMs for
     electrocatalytic purposes. The homogeneous oxidation and the
     electrocatalytic reduction of nitrite and the electrocatalytic oxidation
     of NAD(P)H by an appropriate selection of these V-substituted
     anions were studied.
     85585-38-2
     RL: CAT (Catalyst use); USES (Uses)
        (vanadium-substituted Dawson-type polyoxometalates as
        versatile electrocatalysts)
     85585-38-2 HCAPLUS
ΡN
     Vanadate(8-), [nonacosa-μ-oxohexadecaoxo[μ9-[phosphato(3-)-
CN
     κ0:κ0:κ0:κ0':κ0':κ0'':κ0
     '':\kappa0''':\kappa0''']]hexadecatungstate]hepta-\mu-
     oxodioxo[μ9-[phosphato(3-)-κ0:κ0:κ0:κ0'
     :κ0':κ0'':κ0'':κ0''':κ0''']]di-
     (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     72-2 (Electrochemistry)
     Section cross-reference(s): 78, 67
СТ
     polyoxometalates Dawson structure vanadium substituted
     electrocatalysts nitrite redn oxidn; electrocatalytic oxidn NADPH
IT
     Catalysts
        (electrocatalysts; vanadium-substituted Dawson-type
        polyoxometalates as versatile electrocatalysts)
TT
     Cyclic voltammetry
        (of glassy carbon electrode modified with vanadium-substituted
        Dawson-type polyoxometalates in acidic media)
     Reduction potential
IT
        (of vanadium in vanadium-substituted Dawson-type
        polyoxometalates in acidic media)
TΨ
     UV and visible spectra
        (of vanadium-substituted Dawson-type polyoxometalates
        in buffer solns. containing NaNO2 during electrochem. scanning)
TΤ
     Heteropoly acids
     RL: CAT (Catalyst use); USES (Uses)
        (vanadium-substituted Dawson-type polyoxometalates as
        versatile electrocatalysts)
     85585-38-2
IT
     RL: CAT (Catalyst use); USES (Uses)
        (vanadium-substituted Dawson-type polyoxometalates as
        versatile electrocatalysts)
TT
     53-57-6, NAD(P)H
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); RCT (Reactant); PROC (Process); RACT (Reactant or
        (vanadium-substituted Dawson-type polyoxometalates as
        versatile electrocatalysts for oxidation of)
REFERENCE COUNT:
                                THERE ARE 52 CITED REFERENCES AVAILABLE
                         52
                                FOR THIS RECORD. ALL CITATIONS AVAILABLE
                                IN THE RE FORMAT
                     HCAPLUS COPYRIGHT 2006 ACS on STN
L114 ANSWER 4 OF 64
ACCESSION NUMBER:
                         2005:336064 HCAPLUS
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Les Henderson Page 12 571-272-2538

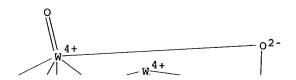
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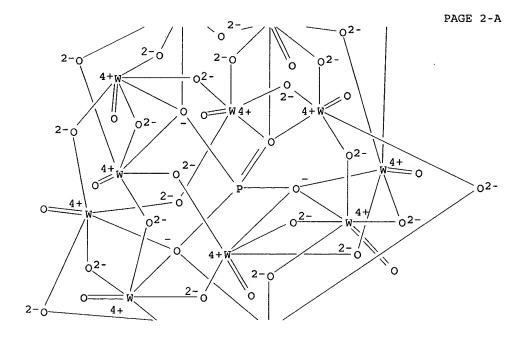
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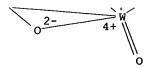
TITLE:

```
Proton-conducting membranes with high
                          selectivity from phosphotungstic acid-
                          doped poly(vinyl alcohol) for DMFC
                          applications
                          Lin, C. W.; Thangamuthu, R.; Yang, C. J.
AUTHOR(S):
CORPORATE SOURCE:
                          Department of Chemical Engineering, National
                          Yunlin University of Science and Technology,
                          Yunlin, 640, Taiwan
SOURCE:
                          Journal of Membrane Science (2005), 253(1-2),
                          23-31
                          CODEN: JMESDO; ISSN: 0376-7388
PUBLISHER:
                          Elsevier B.V.
DOCUMENT TYPE:
                          Journal
LANGUAGE:
                          English
    Proton-conducting hybrid membranes composed of poly(vinyl alc.)
ΔR
     (PVA) and phosphotungstic acid (PWA) were prepared by solution-
     blending. The effect of PWA doping on the
     membrane properties such as water uptake, ion-exchange capacity
     (IEC), proton conductivity and methanol permeability was studied.
     20 weight % PWA content, both water uptake and methanol permeability slightly increased then decreased continuously until 90%. FTIR
     spectra indicate that a significant amount of PWA was maintained in
     the polymer matrix even after several hours of immersion in water.
     Combining FTIR results and the similarity in water uptake and
     methanol permeability of PVA/PWA hybrid membranes suggests that
     the variations in associated properties are ascribed to intermol.
     hydrogen bonding interaction between hydroxyl groups of PVA and
     PWA. Methanol permeabilities of hybrid membranes were
     significantly lower than that of Nafion 115. The proton conductivity of
     hybrid membranes was in the order of 10-4 S cm-1 and generally
     increases with PWA content. Finally, the possibility of PVA/PWA hybrid membranes for use in PEMFC was explored. In
     H2/O2 mode, the performance increases with PWA content and a maximum
     c.d. of 46 mA cm-2 was obtained with PVA 20% and PWA 80%
     (PVA20PWA80) sample. Performance of the DMFC with PVA20PWA80
     increases with temperature and reached to 80 mA cm-2 at 80 °C.
     1343-93-7, Phosphotungstic acid
IT
     RL: DEV (Device component use); PEP (Physical, engineering or
     chemical process); PRP (Properties); PYP (Physical process); PROC
     (Process); USES (Uses)
        (composite blends with polyvinyl alc.;
        proton-conducting membranes with high selectivity from
        phosphotungstic acid-doped poly(vinyl alc.) for DMFC
        applications)
     1343-93-7 HCAPLUS
RN
     Tungstate (3-), tetracosa-\mu-oxododecaoxo [\mu12-[phosphato(3-)-
CN
     κ0:κ0:κ0:κ0':κ0':κ0':κ0'
     ':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec
     a-, trihydrogen (9CI) (CA INDEX NAME)
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PAGE 1-A







●3 H⁴

52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC Section cross-reference(s): 38, 76 proton conducting membrane methanol permeability phosphotungstic ST acid doped; poly vinyl alc heteropoly acid doped fuel cell membrane Cation exchange (PWA acid content effect on; proton-conducting membranes with high selectivity from phosphotungstic acid-doped poly(vinyl alc.) for DMFC applications) ፐጥ Carbon fibers, uses RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses) (cloth, Teflon-coated fibers; proton-conducting membranes with high selectivity from phosphotungstic acid-doped poly(vinyl alc.) for DMFC applications) IT Polyoxyalkylenes, uses RL: DEV (Device component use); PRP (Properties); USES (Uses) (composites with heteropolyacids; proton-conducting membranes with high selectivity from phosphotungstic aciddoped poly(vinyl alc.) for DMFC applications) TΤ Membranes, nonbiological (elec. conductive; proton-conducting membranes with high selectivity from phosphotungstic acid-doped poly(vinyl alc.) for DMFC applications) Polyoxyalkylenes, uses IT RL: DEV (Device component use); PRP (Properties); TEM (Technical or engineered material use); USES (Uses) (fluorine- and sulfo-containing, ionomers; proton-conducting membranes with high selectivity from phosphotungstic acid-doped poly(vinyl alc.) for DMFC applications) IT Electric current-potential relationship (of assembled fuel cells; proton-conducting membranes with high selectivity from phosphotungstic acid-doped poly(vinyl alc.) for DMFC applications) TΤ Membranes, nonbiological (permselective; proton-conducting membranes with high selectivity from phosphotungstic acid-doped poly(vinyl alc.) for DMFC applications) Fluoropolymers, uses TΤ RL: DEV (Device component use); PRP (Properties); TEM (Technical or engineered material use); USES (Uses) (polyoxyalkylene-, sulfo-containing, ionomers; proton-conducting membranes with high selectivity from phosphotungstic aciddoped poly(vinyl alc.) for DMFC applications) TТ Ionomers RL: DEV (Device component use); PRP (Properties); TEM (Technical or engineered material use); USES (Uses) (polyoxyalkylenes, fluorine- and sulfo-containing; proton-conducting membranes with high selectivity from phosphotungstic acid-doped poly(vinyl alc.) for DMFC applications)

IT

Electric energy

(power d. of assembled fuel cells; proton-conducting membranes

```
with high selectivity from phosphotungstic acid-doped
        poly(vinyl alc.) for DMFC applications)
IT
     Fuel cells
        (proton exchange membrane; proton-conducting membranes with
        high selectivity from phosphotungstic acid-doped
        poly(vinyl alc.) for DMFC applications)
TΤ
     Cation exchange membranes
     Current density
       Doping
     Hydrogen bond
     Membrane electrodes
        (proton-conducting membranes with high selectivity from
        phosphotungstic acid-doped poly(vinyl alc.) for DMFC
        applications)
IT
     Ionic conductivity
        (proton; proton-conducting membranes with high selectivity from
        phosphotungstic acid-doped poly(vinyl alc.) for DMFC
        applications)
TΤ
     Permeability
        (to methanol; proton-conducting membranes with high selectivity
        from phosphotungstic acid-doped poly(vinyl alc.) for
        DMFC applications)
     Swelling, physical
ΙT
        (with water; proton-conducting membranes with high selectivity
        from phosphotungstic acid-doped poly(vinyl alc.) for
        DMFC applications)
TТ
     1343-93-7, Phosphotungstic acid
     RL: DEV (Device component use); PEP (Physical, engineering or
     chemical process); PRP (Properties); PYP (Physical process); PROC
     (Process); USES (Uses)
        (composite blends with polyvinyl alc.;
        proton-conducting membranes with high selectivity from
        phosphotungstic acid-doped poly(vinyl alc.) for DMFC
        applications)
     9003-20-7D, Poly(vinyl acetate), 99% hydrolyzed
IT
     RL: DEV (Device component use); PEP (Physical, engineering or
     chemical process); PRP (Properties); PYP (Physical process); PROC
     (Process); USES (Uses)
        (d.p. 2020-2224, composite blends with
        phosphotungstic acid; proton-conducting membranes with high
        selectivity from phosphotungstic acid-doped
        poly(vinyl alc.) for DMFC applications)
TΥ
     67-56-1, Methanol, uses
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PRP (Properties); TEM (Technical or engineered material
     use); PROC (Process); USES (Uses)
        (proton-conducting membranes with high selectivity from
        phosphotungstic acid-doped poly(vinyl alc.) for DMFC
        applications)
TT
     77950-55-1, Nafion 115
     RL: DEV (Device component use); PRP (Properties); TEM (Technical
     or engineered material use); USES (Uses)
        (proton-conducting membranes with high selectivity from
        phosphotungstic acid-doped poly(vinyl alc.) for DMFC
        applications)
     7440-06-4, Platinum, uses
                                 7440-44-0, Carbon, uses
    RL: DEV (Device component use); TEM (Technical or engineered
    material use); USES (Uses)
        (proton-conducting membranes with high selectivity from
        phosphotungstic acid-doped poly(vinyl alc.) for DMFC
        applications)
     7732-18-5, Water, processes
IT
    RL: PEP (Physical, engineering or chemical process); PYP (Physical
    process); PROC (Process)
        (proton-conducting membranes with high selectivity from
        phosphotungstic acid-doped poly(vinyl alc.) for DMFC
```

applications)

REFERENCE COUNT: THERE ARE 30 CITED REFERENCES AVAILABLE 30

FOR THIS RECORD. ALL CITATIONS AVAILABLE

IN THE RE FORMAT

L114 ANSWER 5 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:250678 HCAPLUS

DOCUMENT NUMBER: 142:437700

TITLE: Effective charge transport in

poly(3,4-ethylenedioxythiophene) based hybrid

films containing polyoxometalate

redox centers

AUTHOR (S): Adamczyk, Lidia; Kulesza, Pawel J.;

Miecznikowski, Krzysztof; Palys, Barbara;

Chojak, Malgorzata; Krawczyk, Dorota

Department of Materials and Process CORPORATE SOURCE:

Engineering and Applied Physics, Czestochowa

University of Technology, Czestochowa,

PL-42-200, Pol. Journal of the Electrochemical Society (2005), SOURCE:

152(3), E98-E103

CODEN: JESOAN; ISSN: 0013-4651

PUBLISHER: Electrochemical Society

Journal DOCUMENT TYPE: LANGUAGE: English

Electrodeposition and electrochem. charging of hybrid organic/inorg. films composed of the poly(3,4-ethylenedioxythiophene), PEDOT, conducting polymer matrix, and Keggin type polyoxometalate , phosphododecamolybdate (PMo120403-) or phosphododecatungstate (PW12O403-), redox centers, are described under conditions of aqueous solns. The systems are electropolymd. through potential cycling as thin and moderately thick (μm level) films on electrode surfaces. They are capable of fast charge propagation during redox reactions in strong acid medium (0.5 mol dm-3 H2SO4). high overall physicochem. stability of PEDOT is explored to produce a robust, conductive, matrix for such polynuclear mixed-valence inorg. nanostructures as PMo120403- and PW12O403-. The composite (hybrid) materials are stabilized due to the existence of electrostatic attraction between anionic phosphomolybdate or phosphotungstate units and pos. charged conducting polymer (oxidized). Charge transport is facilitated by the fact that the reversible and fast redox reactions of polyoxometalate appear in the potential range where PEDOT is conductive. The effective diffusion coeffs. are on the level 4 + 10-8 cm2 s-1. The whole concept may lead to the fabrication of composite (hybrid) films that are capable of effective accumulation and propagation of charge in redox capacitors.

12379-13-4 12534-77-9

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)

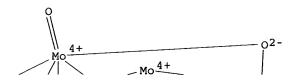
(cyclic voltammetry of poly(ethylenedioxythiophene) containing molybdophosphate or tungstophosphate in H2SO4 solution effective charge transport in hybrid films)

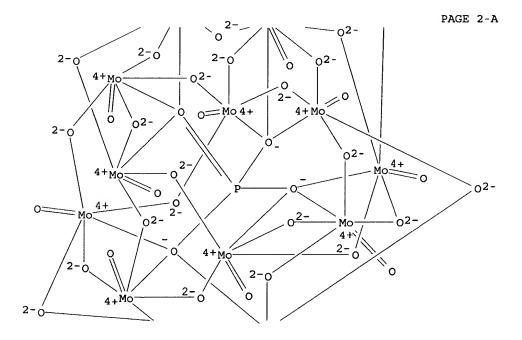
RN 12379-13-4 HCAPLUS

Molybdate (3-), tetracosa- μ -oxododecaoxo [μ 12-[phosphato(3-)-CN INDEX NAME)

Les Henderson 571-272-2538 Page 17

PAGE 1-A

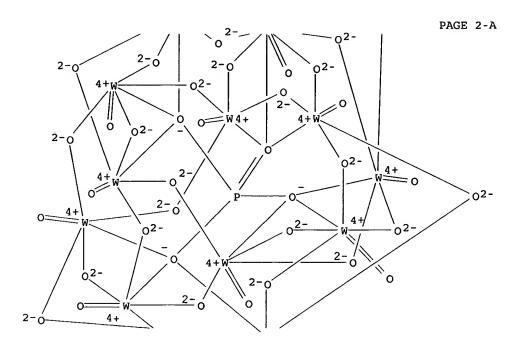




RN 12534-77-9 HCAPLUS CN

Tungstate(3-), tetracosa-μ-oxododecaoxo[μ12-[phosphato(3-)-0:0:0':0':0':0'':0'':0''':0''']]dodeca-(9CI) (CA INDEX NAME)

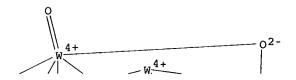


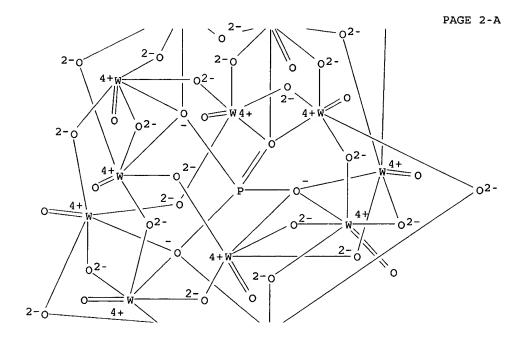


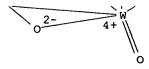


1343-93-7, Tungstophosphoric acid H3PW12O40 ΙT 12026-57-2, Molybdophosphoric acid h3pmo12o40 RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process) (electrochem. polymerization of ethylenedioxythiophene in aqueous solution containing H3PMo12040 and H3PW12040 and effective charge transport in poly(ethylenedioxythiophene) based hybrid films containing polyoxometalate redox centers) RN 1343-93-7 HCAPLUS Tungstate (3-), tetracosa- μ -oxododecaoxo [μ 12-[phosphato(3-)-CN κ0:κ0:κ0:κ0':κ0':κ0':κ0' ':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec a-, trihydrogen (9CI) (CA INDEX NAME)

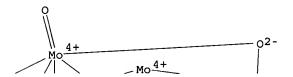
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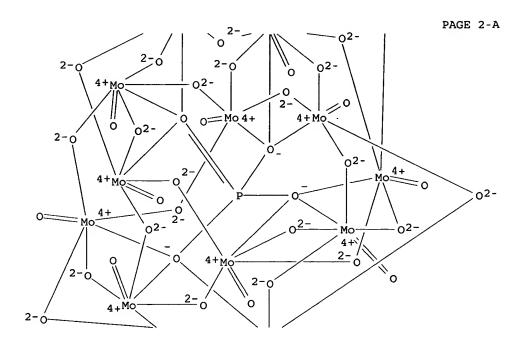


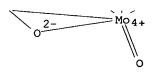


●3 H+



571-272-2538





PAGE 3-A

●3 H+

CC 72-2 (Electrochemistry)

Section cross-reference(s): 35, 36, 67, 78

ST effective charge transport polyethylenedioxythiophene hybrid film polyoxometalate center; electropolymn

ethylenedioxythiophene molybdophosphoric tungstophosphoric acid

IT Polymerization

(electrochem.; of ethylenedioxythiophene in aqueous solution containing H3PMo12O4O and H3PW12O4O and effective charge transport in poly(ethylenedioxythiophene) based hybrid films containing polyoxometalate redox centers)

IT Redox reaction

(electrochem.; of molybdophosphate or tungstophosphate in poly(ethylenedioxythiophene) films in H2SO4 solution and effective charge transport in poly(ethylenedioxythiophene) based hybrid films containing polyoxometalate redox centers)

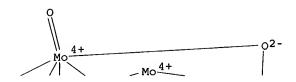
IT Reduction, electrochemical

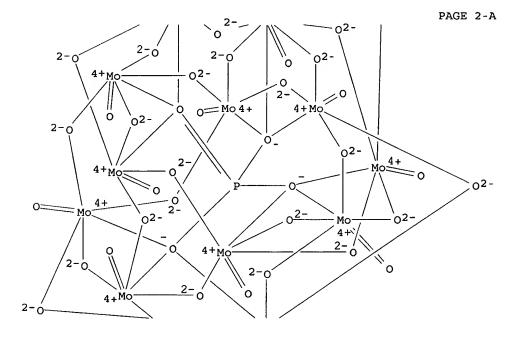
(of molybdophosphate or tungstophosphate in poly(ethylenedioxythiophene) films in H2SO4 solution and effective charge transport in poly(ethylenedioxythiophene) based hybrid films containing polyoxometalate redox centers)

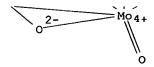
IT 12379-13-4 12534-77-9

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)

```
(cyclic voltammetry of poly(ethylenedioxythiophene) containing
        molybdophosphate or tungstophosphate in H2SO4 solution effective
        charge transport in hybrid films)
     126213-50-1, 3,4-Ethylenedioxythiophene
TТ
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (electrochem. polymerization in aqueous solution containing molybdophosphoric
        acid or tungstophosphoric acid and effective charge transport
        in poly(ethylenedioxythiophene) based hybrid films containing
        polyoxometalate redox centers)
     1343-93-7, Tungstophosphoric acid H3PW12O40
TΤ
     12026-57-2, Molybdophosphoric acid h3pmo12o40
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PROC (Process)
        (electrochem. polymerization of ethylenedioxythiophene in aqueous solution
        containing H3PMo12040 and H3PW12040 and effective charge transport
        in poly(ethylenedioxythiophene) based hybrid films containing
        polyoxometalate redox centers)
REFERENCE COUNT:
                         61
                                THERE ARE 61 CITED REFERENCES AVAILABLE
                                FOR THIS RECORD. ALL CITATIONS AVAILABLE
                                IN THE RE FORMAT
L114 ANSWER 6 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2005:102386 HCAPLUS
DOCUMENT NUMBER:
                          142:338886
TITLE:
                         Preferential oxidation of CO in H2 by aqueous
                         polyoxometalates over metal catalysts
                         Kim, Won Bae; Voitl, Tobias; Rodriguez-Rivera,
AUTHOR(S):
                         Gabriel J.; Evans, Steven T.; Dumesic, James
CORPORATE SOURCE:
                         Chemical and Biological Engineering
                         Department, University of Wisconsin, Madison,
                         WI, 53706, USA
SOURCE:
                         Angewandte Chemie, International Edition
                          (2005), 44(5), 778-782
                         CODEN: ACIEF5; ISSN: 1433-7851
                         Wiley-VCH Verlag GmbH & Co. KGaA
PUBLISHER:
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     Stream cleaning: CO in CO/H2 mixts. is
     oxidized preferentially at room temperature with an aqueous
     polyoxometalate (POM) solution over gold catalysts. The rate
     of H2 oxidation is slow and is inhibited by CO. This process can be
     used to remove CO efficiently from H2 gas streams. The solution
     containing protons and reduced POM can be used to produce elec. energy
     at a fuel-cell anode through reoxidn. of the reduced POM.
     12026-57-2, Dodecamolybdophosphoric acid
TΤ
     RL: FMU (Formation, unclassified); RCT (Reactant); FORM (Formation, nonpreparative); RACT (Reactant or reagent)
        (preferential oxidation of CO in H2 by aqueous
        polyoxometalates over metal catalysts)
RN
     12026-57-2 HCAPLUS
     Molybdate(3-), tetracosa-\mu-oxododecaoxo[\mu12-[phosphato(3-)-
CN
     κ0:κ0:κ0:κ0':κ0':κ0':κ0'
     ':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec
     a-, trihydrogen (9CI) (CA INDEX NAME)
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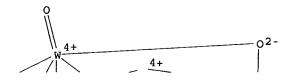


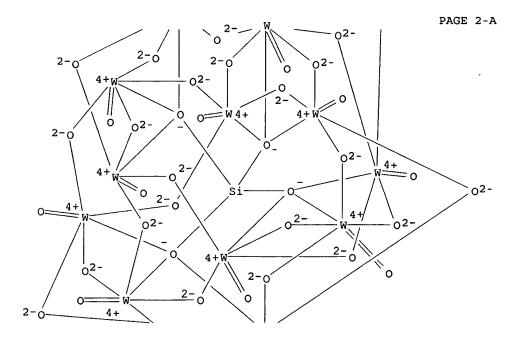


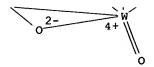
●3 H⁺

52-1 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 67 ST preferential oxidn carbon monoxide polyoxometalate metal heterogeneous catalysis; fuel cell stream cleanup selective oxidn catalyst heteropoly acid TT Nanoparticles (catalyst; preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) IT Reduction (of polyoxometalate; preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) IT Fuel cell anodes (preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) TТ Heteropoly acids RL: CPS (Chemical process); FMU (Formation, unclassified); PEP (Physical, engineering or chemical process); FORM (Formation, nonpreparative); PROC (Process) (preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) IT Oxidation, electrochemical (reoxidn. of polyoxometalate; preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) TT Oxidation Oxidation catalysts Oxidation kinetics (selective; preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) ፐጥ 7440-44-0, Carbon, uses RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES (Uses) (catalyst support; preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) TΤ 7440-57-5, Gold, uses RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES (Uses) (nanotubes; preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) 7440-15-5P, Rhenium, uses 7440-17-7P, Rubidium, uses 7440-22-4P, Silver, uses RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses) (preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) IT. 7439-88-5, Iridium, uses 7440-05-3, Palladium, uses 7440-06-4, Platinum, uses RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES (Uses) (preferential oxidation of CO in H2 by aqueous polyoxometalates over metal catalysts) 630-08-0, Carbon monoxide, processes RL: CPS (Chemical process); PEP (Physical, engineering or chemical IT

```
process); REM (Removal or disposal); PROC (Process)
        (preferential oxidation of CO in H2 by aqueous
        polyoxometalates over metal catalysts)
     12026-57-2, Dodecamolybdophosphoric acid
TT
     RL: FMU (Formation, unclassified); RCT (Reactant); FORM
     (Formation, nonpreparative); RACT (Reactant or reagent)
        (preferential oxidation of CO in H2 by aqueous
        polyoxometalates over metal catalysts)
IT
     1333-74-0, Hydrogen, reactions
     RL: OCU (Occurrence, unclassified); RCT (Reactant); OCCU
     (Occurrence); RACT (Reactant or reagent)
        (preferential oxidation of CO in H2 by aqueous
        polyoxometalates over metal catalysts)
                                THERE ARE 25 CITED REFERENCES AVAILABLE
REFERENCE COUNT:
                          25
                                FOR THIS RECORD. ALL CITATIONS AVAILABLE
                                IN THE RE FORMAT
L114 ANSWER 7 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                          2005:15474 HCAPLUS
DOCUMENT NUMBER:
                          142:359869
TITLE:
                          Screening of polyoxometalates,
                          semiconductor metal oxides and zeolites for
                          photocatalytic activity and selectivity
AUTHOR (S):
                          Arslan-Alaton, Idil; Selcuk, Huseyin;
                          Erdem-Senatalar, Ayse
Faculty of Civil Engineering, Department of
CORPORATE SOURCE:
                          Environmental Engineering, Istanbul Technical
                          University, Istanbul, 34469, Turk.
SOURCE:
                          Fresenius Environmental Bulletin (2004),
                          13(11b, y), 1248-1252
CODEN: FENBEL; ISSN: 1018-4619
                          PSP - Parlar Scientific Publications
PUBLISHER:
                          Journal
DOCUMENT TYPE:
LANGUAGE:
                          English
     This study reports the treatment performance and selectivity of 2
     photochem. active materials, namely TiO2 (1 g/L, pH 2.8) and the
     heteropolyacid H4SiW12O40 (1.44 g/L or 0.5mM in aqueous solution, pH 2.8)
     for the oxidation of phenol, iso-Pr alc. and formic acid selected as
     the recalcitrant index pollutants, using UV-C (20 W) and
     UV-A (125 W) light sources in 2 different photoreactors. In sep.
     expts., the effect of the zeolites Y (1 g/L; Si/Al =5.2/1 and
     80/1) and Beta (1 g/L; Si/Al =75/1) as TiO2 and H4SiW12O40
     catalyst supports was studied under the same reaction conditions.
     It could be demonstrated that H4SiW12O40 is more selective than
     TiO2, especially for the charge transfer type isopropanol oxidation It was
     also found that the zeolitic supports improved the photocatalytic
     activity of polyoxometalates (POM) with a pronounced
     effect for UV-C (short-UV) irradiation A significantly reduced photoactivity (12.5% for UV-A and 9.5% for UV-C) in comparison
     with TiO2-mediated photocatalysis was observed for the
     combination POM + TiO2, speculatively due to the
     conduction band electron short-circuit effect of POM, i.e. ecb +
     POM → POMred.
     12027-38-2
IT
     RL: CAT (Catalyst use); USES (Uses)
        (screening of polyoxometalates and semiconductor
        metal oxides and zeolites for photocatalytic activity and
        selectivity)
RN
     12027-38-2 HCAPLUS
     Tungstate(4-), [\mu 12-[orthosilicato(4-)-
CN
     κ0:κ0:κ0:κ0':κ0':κ0':κ0'
     ':κ0'':κ0'':κ0''':κ0''':κ0''']]tetra
     cosa-µ-oxododecaoxododeca-, tetrahydrogen (9CI) (CA INDEX
     NAME)
```







●4 H+

60-2 (Waste Treatment and Disposal) Section cross-reference(s): 67 ST polyoxometalate semiconductor metal oxide zeolite photocatalysis IT Wastewater treatment (photocatalytic; screening of polyoxometalates and semiconductor metal oxides and zeolites for photocatalytic activity and selectivity) Catalysis Catalysts (photochem.; screening of polyoxometalates and semiconductor metal oxides and zeolites for photocatalytic activity and selectivity) IT Catalyst supports (screening of polyoxometalates and semiconductor metal oxides and zeolites for photocatalytic activity and selectivity) IT H-Beta zeolites Heteropoly acids Oxides (inorganic), uses Y zeolites Zeolite NaY Zeolites (synthetic), uses RL: CAT (Catalyst use); USES (Uses) (screening of polyoxometalates and semiconductor metal oxides and zeolites for photocatalytic activity and selectivity) 13463-67-7, Titania, uses IT 12027-38-2 RL: CAT (Catalyst use); USES (Uses) (screening of polyoxometalates and semiconductor metal oxides and zeolites for photocatalytic activity and selectivity) 64-18-6, Formic acid, processes 67-63 processes 108-95-2, Phenol, processes ΙT 67-63-0, Iso-propyl alcohol, RL: REM (Removal or disposal); PROC (Process) (screening of polyoxometalates and semiconductor metal oxides and zeolites for photocatalytic activity and selectivity) REFERENCE COUNT: 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L114 ANSWER 8 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2005:5417 HCAPLUS DOCUMENT NUMBER: 142:359724 TITLE: Method for desulfurizing and denitrifying flue gas simultaneously INVENTOR(S): Wang, Rui PATENT ASSIGNEE(S): Hainan University, Peop. Rep. China SOURCE: Faming Zhuanli Shenqing Gongkai Shuomingshu, 5

CODEN: CNXXEV

Patent

DOCUMENT TYPE:

LANGUAGE:

Chinese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 1478588	Α	20040303	CN 2003-147304	
				2003
				0704
PRIORITY APPLN. INFO.:			CN 2003-147304	
				2003
				0704

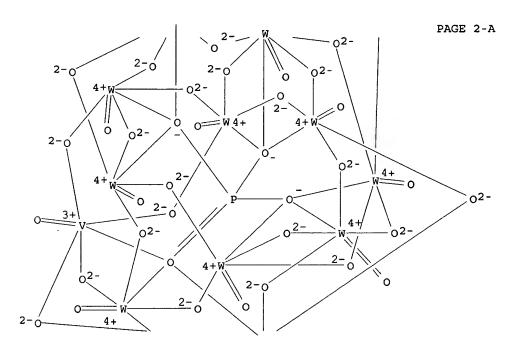
The absorbent of heteropoly acid or its reduced form for desulfurizing and denitrifying flue gas simultaneously is prepared from phosphate or silicate, molybdate, and tungstate. The additive (such as NaCl, CuSO4, H3PO4, ZnSO4, MnSO4, CuS, and/or NH4VO3) may be mixed with the absorbent . IT 12398-73-1P

RL: CPS (Chemical process); NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process); USES (Uses) (method for desulfurizing and denitrifying flue gas simultaneously using heteropoly acid absorbent)

RN

12398-73-1 HCAPLUS
Vanadate(4-), (eicosa-µ-oxoundecaoxoundecatungstate)tetra-µ-CN $oxooxo[\mu12-[phosphato(3-)-\kappa0:\kappa0:\kappa0:\kappa0]$: κ0':κ0':κ0'':κ0'':κ0'':κ0''':. kappa.O''':κO''']]-, tetrahydrogen (9CI) (CA INDEX NAME)

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT



0 2- W

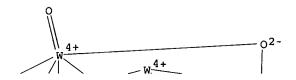
PAGE 3-A

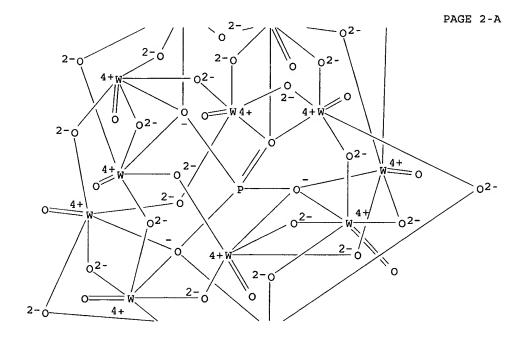
●4 H+

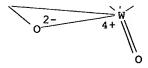
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ICM B01D053-78
TC
     ICS B01D053-60
CC
     59-4 (Air Pollution and Industrial Hygiene)
     12398-73-1P 63950-64-1P 477978-49-7P RL: CPS (Chemical process); NUU (Other use, unclassified); PEP
     (Physical, engineering or chemical process); SPN (Synthetic
     preparation); PREP (Preparation); PROC (Process); USES (Uses)
        (method for desulfurizing and denitrifying flue gas
        simultaneously using heteropoly acid absorbent)
L114 ANSWER 9 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                          2004:1119124 HCAPLUS
DOCUMENT NUMBER:
                          142:231812
                          Synthesis, optical properties and electronic
TITLE:
                          structures of polyoxometalates
                          K3P(Mo1-xWx)12040 (0 \le x \le 1)
AUTHOR (S):
                          Goubin, F.; Guenee, L.; Deniard, P.; Koo,
                          H.-J.; Whangbo, M.-H.; Montardi, Y.; Jobic, S.
CORPORATE SOURCE:
                          Institut des Materiaux Jean Rouxel,
                          Laboratoire de Chimie des Solides, BP 32229,
                          UMR 6502 CNRS-Universite de Nantes, Nantes,
                          44322, Fr.
SOURCE:
                          Journal of Solid State Chemistry (2004),
                          177(12), 4528-4534
                          CODEN: JSSCBI; ISSN: 0022-4596
PUBLISHER:
                          Elsevier
DOCUMENT TYPE:
                          Journal
LANGUAGE:
                          English
     Various compns. of solid solns. K3P (Mo1-xWx) 12040
     (0≤x≤1) were prepared using two solid state synthetic
     routes. The crystallite size was determined by linewidth refinements
     of x-ray diffraction patterns using the Warren-Averbach
     method, and the grain size distribution by laser scattering expts.
     Optical properties were determined by diffuse reflectance measurements
     in the UV-visible range. The optical gap Eq increases
     exponentially from .apprx.2.5 to .apprx.3.30 eV with increasing x,
     and is systematically shifted to a higher energy when the grain
     size decreases. The relation between Eq and x was analyzed by
     calculating the HOMO-LUMO gaps of the [P(Mo1-xWx)12040]3- anions from
     tight-binding electronic structure calcns.
TT
     1343-93-7, 12-Tungstophosphoric acid 12026-57-2,
     12-Molybdophosphoric acid
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (for preparation of solid solns. of molybdophosphates with
        tungstophosphates)
RN
     1343-93-7 HCAPLUS
CN
     Tungstate(3-), tetracosa-\mu-oxododecaoxo[\mu12-[phosphato(3-)-
     κ0:κ0:κ0:κ0':κ0':κ0':κ0'
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':κΟ'':κΟ'':κΟ''':κΟ''']]dodec a-, trihydrogen (9CI) (CA INDEX NAME)

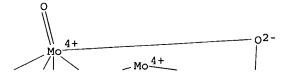
PAGE 1-A

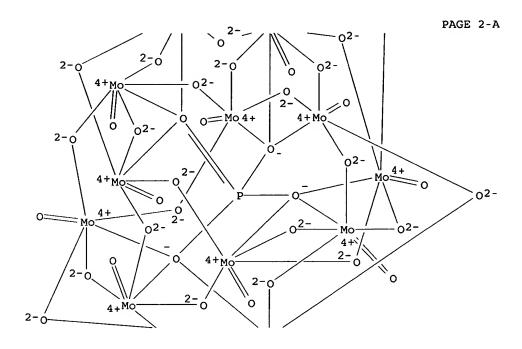


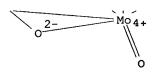




●3 H⁴







●3 H+

IT 12026-68-5P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(preparation via acid route vs. oxide route, particle size distribution and reflectance spectrum of)

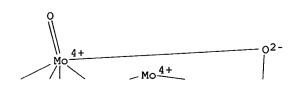
RN

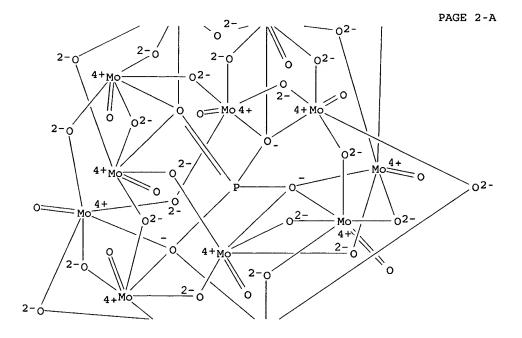
12026-68-5 HCAPLUS
Molybdate(3-), tetracosa-μ-oxododecaoxo[μ12-[phosphato(3-)-CN

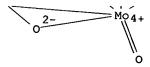
a-, tripotassium (9CI) (CA INDEX NAME)

Les Henderson Page 34 571-272-2538

PAGE 1-A

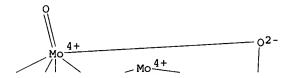


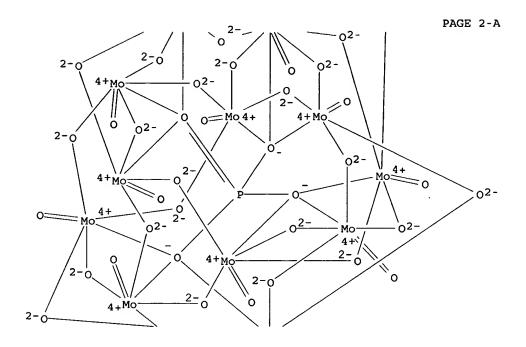


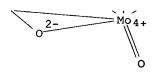


■3 K⁴

PAGE 1-A

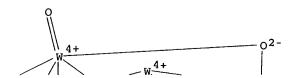


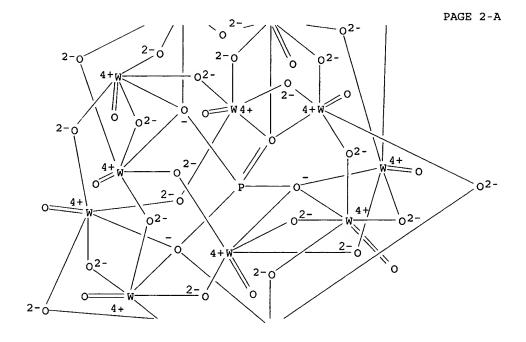




●3 K+

PAGE 1-A





02- W

PAGE 3-A

●3 K⁺

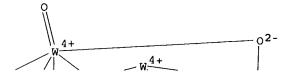
78-7 (Inorganic Chemicals and Reactions) CC Section cross-reference(s): 75 TT 584-08-7, Potassium carbonate (K2CO3) 1313-27-5, Molybdenum oxide (MoO3), reactions 1314-35-8, Tungsten oxide (WO3), reactions 1343-93-7, 12-Tungstophosphoric acid 7758-11-4, Potassium phosphate (K2HPO4) 12026-57-2, 12-Molybdophosphoric acid RL: RCT (Reactant); RACT (Reactant or reagent) (for preparation of solid solns. of molybdophosphates with tungstophosphates) IT 12026-68-5P RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (preparation via acid route vs. oxide route, particle size distribution and reflectance spectrum of) 12026-68-5DP, solid solns. with tungsten analog 12207-66-8P RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (preparation, crystal structure, optical properties and electronic structures of) REFERENCE COUNT: THERE ARE 28 CITED REFERENCES AVAILABLE 28 FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L114 ANSWER 10 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2004:1068364 HCAPLUS DOCUMENT NUMBER: 142:284416 TITLE: Efficient degradation of dye pollutants on nanoporous polyoxotungstate-anatase composite under visible-light irradiation AUTHOR (S): Yang, Yu; Wu, Qingyin; Guo, Yihang; Hu, Changwen; Wang, Enbo CORPORATE SOURCE: Faculty of Chemistry, Northeast Normal University, Changchun, 130024, Peop. Rep. China SOURCE: Journal of Molecular Catalysis A: Chemical (2005), 225(2), 203-212 CODEN: JMCCF2; ISSN: 1381-1169 PUBLISHER: Elsevier B.V. DOCUMENT TYPE: Journal English LANGUAGE: A novel photocatalyst, nanoporous anatase TiO2 crystalline particles coupled by homogeneously dispersed Keggin unit, H3PW12O4O/TiO2, was prepared by a simple and rapid process, i.e., at a lower temperature (200°) by combined sol-gel and programmed temperature hydrothermal methods. The resulting material was characterized by UV diffuse reflectant spectroscopy, XRD, 31P MAS NMR, TEM, and N

adsorption. This new photocatalyst exhibited visible-light

photocatalytic activity to decompose 10 various organic dyes in aqueous

systems. It was attempted to determine the feasibility of such a degradation by varying the chemical structures, either azoic (Congo Red, Methyl orange, Ponceau G, Orange II, and Eriochrome Blue Black B), or anthraquinonic (Alizarin S), or heteropolyarom. (Methylene Blue), or fluorescent (Neutral Red, Rhodamine B), or sulfonic (Fuchsin Acid). The intermediates and the final products of degradation were detected by electrospray mass spectrometer and ion chromatog. According to the exptl. results, we proposed a possible mechanism of the photodegrdn. of dyes under visible-light irradiation in the aqueous system. 1343-93-7DP, Tungstophosphoric acid (H3PW12O40), titania coupled with RL: CAT (Catalyst use); PRP (Properties); SPN (Synthetic preparation); PREP (Preparation); USES (Uses) (preparation and characterization of nanoporous anatase TiO2 coupled by H3PW12O40 and its activity as photocatalyst for degradation of dyes in aqueous systems under visible-light irradiation) 1343-93-7 HCAPLUS Tungstate(3-), tetracosa-μ-oxododecaoxo[μ12-[phosphato(3-)κ0:κ0:κ0:κ0':κ0':κ0':κ0' ':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec

PAGE 1-A

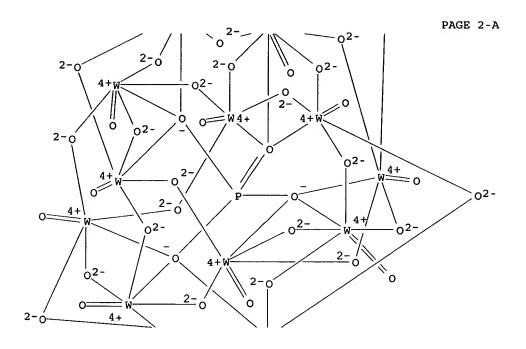


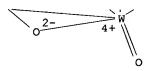
a-, trihydrogen (9CI) (CA INDEX NAME)

IT

RN

CN





●3 н+

CC 60-2 (Waste Treatment and Disposal)

Section cross-reference(s): 22, 41, 67, 74

ST photocatalyst nanoporous anatase titania polyoxotungstate composite prepn; wastewater dye photodegrdn catalyst

polyoxotungstate anatase nanoporous composite

1343-93-7DP, Tungstophosphoric acid (H3PW12O40), titania coupled with 13463-67-7DP, Titania, coupled by H3PW12O40 RL: CAT (Catalyst use); PRP (Properties); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(preparation and characterization of nanoporous anatase TiO2 coupled by H3PW12O4O and its activity as photocatalyst for degradation of dyes in aqueous systems under visible-light irradiation)

REFERENCE COUNT:

44 THERE ARE 44 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L114 ANSWER 11 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2004:1022769 HCAPLUS

DOCUMENT NUMBER: 143:288170

TITLE: Use of an interactive carbohydrate polymer

phenomenon for the separation of

polyoxometalates

AUTHOR(S): Ruuttunen, Kyoesti; Vuorinen, Tapani CORPORATE SOURCE: Laboratory of Forest Products Chemistry,

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Helsinki University of Technology, Espoo, FIN-02015, Finland Carbohydrate Polymers (2004), 58(4), 443-448
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SOURCE: Carbohydrate Polymers (2004), 58(4), 443
CODEN: CAPOD8; ISSN: 0144-8617

PUBLISHER: Elsevier B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

AB Keggin type polyoxometalates (POMs) are proposed as agents for improving the feasibility of oxygen bleaching technique used in producing cellulose fiber in the wood pulping industry. To study the possible interactions between carbohydrate polymers and POMs, several different POM anions were eluted through a column containing dextran chromatog. gel (Sephadex G-50). Solns. of KCl (0.05 and 0.10 mol/L) and NaCl (0.10 mol/L) were used as the eluents. The elution vols. (Ve) of all the POMs tested are greater than the Ve of the internal standard, glycine. The Ve of the POMs increased along with increasing electrolyte concentration When eluted with 0.10 mol/L KCl solution, the Ve were greater than the resp. values of the same POMs eluted with 0.10 mol/L NaCl solution These facts suggest that POM and the Sephadex gel form complexes, and that the tendency for this to take place depends on the type and concentration of cations present in the system. The method described in this paper may be useful in analyzing and purifying mixts. of POMs.

IT 93279-93-7 137531-01-2

RL: ANT (Analyte); TEM (Technical or engineered material use);
ANST (Analytical study); USES (Uses)

(separation of polyoxometalates by gel filtration using Sephadex G50 column)

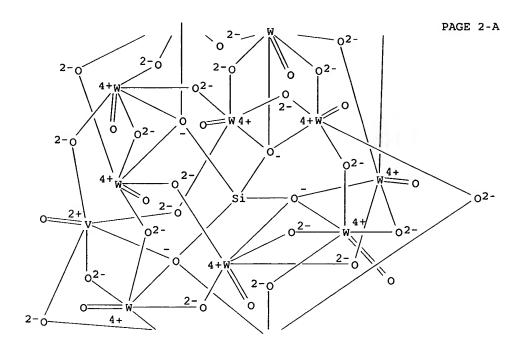
RN 93279-93-7 HCAPLUS

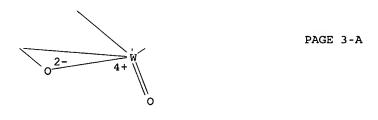
Vanadate(6-), (eicosa-μ-oxoundecaoxoundecatungstate) [μ12[orthosilicato(4-)-κ0:κ0:κ0':κ0':
κ0':κ0'':κ0'':κ0''':κ0'''

:κΟ''']]tetra-μ-οχοοχο-, hexapotassium (9CI) (CA INDEX NAME)

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

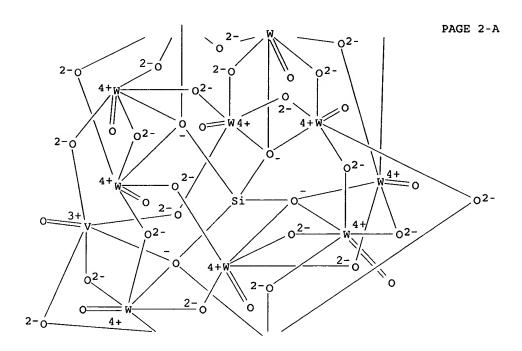
Les Henderson Page 42 571-272-2538

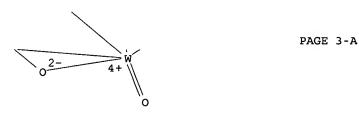




●6 K+

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





●5 K+

CC 43-6 (Cellulose, Lignin, Paper, and Other Wood Products) Section cross-reference(s): 79 ST Keggin polyoxometalate sepn gel filtration Sephadex G50 column; pulp bleaching agent polyoxometalate sepn gel filtration IT Pulp bleaching (oxygen; separation of polyoxometalates by gel filtration using Sephadex G50 column) IT Size-exclusion chromatography (separation of polyoxometalates by gel filtration using Sephadex G50 column) IT Heteropoly acids RL: ANT (Analyte); ANST (Analytical study) (tungstates; separation of polyoxometalates by gel filtration using Sephadex G50 column) IT **93279-93-7 137531-01-2** 172304-20-0 394212-03-4 680592-64-7 864384-24-7 RL: ANT (Analyte); TEM (Technical or engineered material use); ANST (Analytical study); USES (Uses) (separation of polyoxometalates by gel filtration using Sephadex G50 column)

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9048-71-9, Sephadex G 50
TΤ
```

RL: ARU (Analytical role, unclassified); NUU (Other use, unclassified); ANST (Analytical study); USES (Uses)

(separation of polyoxometalates by gel filtration using

Sephadex G50 column)

REFERENCE COUNT:

THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE

IN THE RE FORMAT

L114 ANSWER 12 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2004:999712 HCAPLUS

DOCUMENT NUMBER:

141:427184

TITLE:

Compositions, materials

incorporating the compositions, and methods of using the compositions

and materials

INVENTOR(S):

Okun, Nelya; Hill, Craig L.

PATENT ASSIGNEE(S): USA

SOURCE:

U.S. Pat. Appl. Publ., 8 pp.

CODEN: USXXCO

DOCUMENT TYPE: LANGUAGE:

Patent English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

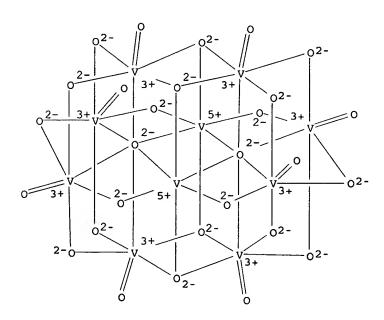
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						-									
US	US 2004230086				A1	:	20041118			US 2004-786671					
													2004 0225		
WO 2005021435				A2	:	20050310		,	WO 2004-US5645					0223	
															2004 0225
	W:						AU,	-		-	-		-	-	-
		-		-			CU,			-					-
					•		GH,		,		•		•	•	•
							LC,		-						•
		-	-	-			MZ,		-						-
				•	•		SE,	•			•	•	•	TN,	TR,
							UΖ,		•		-				
	RW:		•	•			MW,		•	•	•	•		•	•
							MD,							•	•
			-				FI,								-
					•	•	SK,		•		CF,	CG,	CI,	CM,	GA,
		•		•	ML,	MR,	NE,	SN,							
PRIORITY APPLN. INFO.:										US 2003-449892P]	?
															2003 0225
							US 2004-786671					71	1	A	
															2004
															0225

AB Compns. that can protect and/or remove contaminants such as warfare agents from the environment in which people are operating are disclosed, as are materials incorporating the compns., and methods of use thereof. In one embodiment, the composition includes a metal nitrate selected from d-block metal nitrates and f-block metal nitrates and a metal salt having weakly bound counter anions. The metal of the metal salt having weakly bound counter anions is selected from a d-block metal and an f-block metal. Another embodiment of the composition includes a first polyoxometalate having a first metal selected from a d-block metal and an f-block metal and a second

polyoxometalate having a second metal selected from a d-block metal and an f-block metal, the first metal being an open coordinate site of the first polyoxometalate. In addition, the first metal has a nitrate terminal ligand. 59858-44-5 134360-58-0 795308-36-0 IT 796042-78-9 RL: CAT (Catalyst use); USES (Uses) (as polyoxometalate; catalytic compns. for removal of contaminants such as warfare agents, and materials incorporating these compns.) RN 59858-44-5 HCAPLUS 1-Butanaminium, N,N,N-tributyl-, tetradeca-μ-oxo-tetra-μ3-CN oxodi-µ6-oxooctaoxodecavanadate(6-) (6:1) (9CI) (CA INDEX NAME)

CM 1

CRN 12397-12-5 CMF 028 V10 CCI CCS



CM 2

CRN 10549-76-5 CMF C16 H36 N

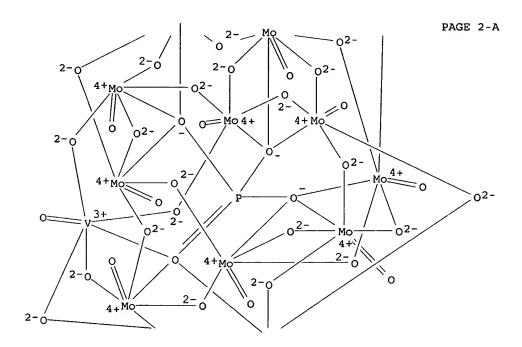
RN 134360-58-0 HCAPLUS

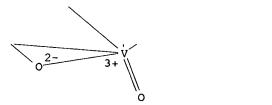
CN 1-Butanaminium, N,N,N-tributyl-, (heptadeca-μoxodecaoxodecamolybdate)hepta-μ-oxodioxo[μ12-[phosphato(3-)κΟ:κΟ:κΟ:κΟ':κΟ':κΟ'

':κΟ'':κΟ'':κΟ''':κΟ''':κΟ''']]divan adate(5-) (5:1) (9CI) (CA INDEX NAME) CM 1

CRN 58071-93-5 CMF Mo10 O40 P V2 CCI CCS

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





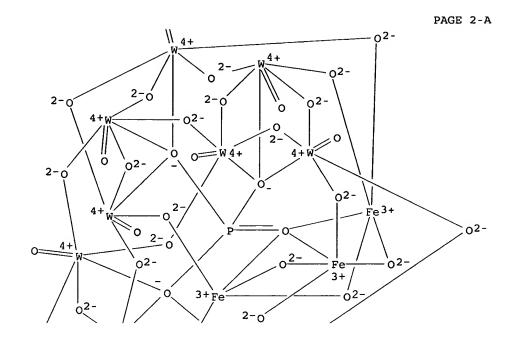
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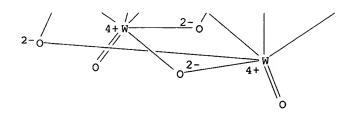
CRN 10549-76-5 CMF C16 H36 N PAGE 3-A

PAGE 1-A

0

CMF Fe3 O37 P W9





CM 2

CRN 10549-76-5 CMF C16 H36 N

CMF Fe3 O68 P2 W18

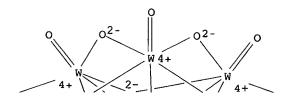
RN 796042-78-9 HCAPLUS
CN 1-Butanaminium, N,N,N-tributyl-, triferratedotetraconta-μoxooctadecaoxobis[μ9-[phosphato(3-)κΟ:κΟ:κΟ:κΟ':κΟ':κΟ'
'':κΟ'':κΟ'':κΟ'
'':κΟ''':κΟ''']]octadecatungstate(9-) (9:1) (9CI) (CA
INDEX NAME)

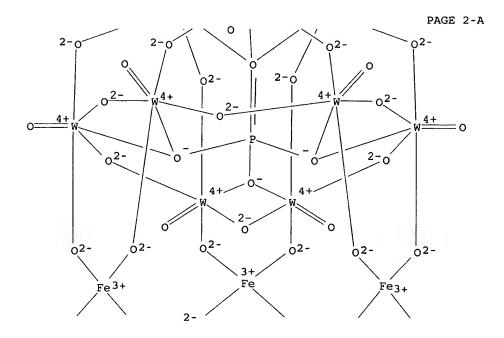
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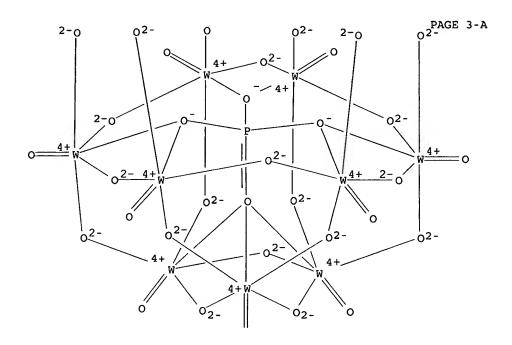
CRN 796042-77-8

CCI CCS

PAGE 1-A







PAGE 4-A

0

CM 2 CRN 10549-76-5 CMF C16 H36 N

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 Cerium (III) nitrate 10141-05-6, Cobalt (II) nitrate
 10421-48-4, Iron (III) nitrate 13093-17-9
 13138-45-9, Nickel (II) nitrate 13770-18-8,
 Copper (II) perchlorate 34946-82-2, Copper (II)
 trifluoromethanesulfonate 38465-60-0, Copper (II)
 tetrafluoroborate
 RL: CAT (Catalyst use); USES (Uses)
 (catalytic compns. for removal of
 contaminants such as warfare agents, and
 materials incorporating these compns.)
RN 3251-23-8 HCAPLUS
CN Nitric acid, copper(2+) salt (8CI, 9CI) (CA INDEX NAME)

●1/2 Cu(II)

RN 10108-73-3 HCAPLUS
CN Nitric acid, cerium(3+) salt (8CI, 9CI) (CA INDEX NAME)

●1/3 Ce(III)

RN 10141-05-6 HCAPLUS CN Nitric acid, cobalt(2+) salt (8CI, 9CI) (CA INDEX NAME)

●1/2 Co(II)

RN 10421-48-4 HCAPLUS CN Nitric acid, iron(3+) salt (8CI, 9CI) (CA INDEX NAME)

●1/3 Fe(III)

RN 13093-17-9 HCAPLUS CN Nitric acid, cerium(4+) salt (8CI, 9CI) (CA INDEX NAME)

●1/4 Ce(IV)

RN 13138-45-9 HCAPLUS CN Nitric acid, nickel(2+) salt (8CI, 9CI) (CA INDEX NAME)

●1/2 Ni(II)

RN 13770-18-8 HCAPLUS CN Perchloric acid, copper(2+) salt (8CI, 9CI) (CA INDEX NAME)

●1/2 Cu(II)

RN 34946-82-2 HCAPLUS
CN Methanesulfonic acid, trifluoro-, copper(2+) salt (9CI) (CA INDEX NAME)

●1/2 Cu(II)

RN 38465-60-0 HCAPLUS CN Borate(1-), tetrafluoro-, copper(2+) (2:1) (9CI) (CA INDEX NAME)

●1/2 Cu(II) 2+

IC ICM A62D003-00 ICS C11D001-00

INCL 588205000

CC 59-2 (Air Pollution and Industrial Hygiene)

Section cross-reference(s): 4

ST polyoxymetalate nitrate copper catalytic oxidn warfare agent

IT Biological warfare agents Chemical warfare agents Coating materials

```
Environmental pollution control
     Oxidation catalysts
     Powders
     Textiles
         (catalytic compns. for removal of
         contaminants such as warfare agents, and
         materials incorporating these compns.)
IT
     Aldehydes, processes
     RL: ADV (Adverse effect, including toxicity); CPS (Chemical process); PEP (Physical, engineering or chemical process); POL
     (Pollutant); REM (Removal or disposal); BIOL (Biological study);
     OCCU (Occurrence); PROC (Process)
         (catalytic compns. for removal of
         contaminants such as warfare agents, and
        materials incorporating these compns.)
TΤ
     Oxidation
         (catalytic; catalytic compns. for removal of
         contaminants such as warfare agents, and
        materials incorporating these compns.)
TΤ
     Drug delivery systems
         (topical; catalytic compns. for removal of
        contaminants such as warfare agents, and
        materials incorporating these compns.)
TΤ
     Heteropoly acids
     RL: CAT (Catalyst use); USES (Uses)
         (tungstates, complexes with iron, silver, and/or cerium;
        catalytic compns. for removal of contaminants
        such as warfare agents, and materials incorporating
         these compns.)
     7727-37-9D, Nitrogen, compds.
TT
     RL: ADV (Adverse effect, including toxicity); CPS (Chemical process); PEP (Physical, engineering or chemical process); POL
     (Pollutant); REM (Removal or disposal); BIOL (Biological study);
     OCCU (Occurrence); PROC (Process)
         (aliphatic; catalytic compns. for removal of
        contaminants such as warfare agents, and
        materials incorporating these compns.)
TТ
     7440-33-7D, Tungsten, heteropoly compds. containing, complexes with
     iron 59858-44-5 134360-58-0
     795308-36-0 796042-78-9
     RL: CAT (Catalyst use); USES (Uses)
         (as polyoxometalate; catalytic compns. for
        removal of contaminants such as warfare
        agents, and materials incorporating these compns.)
IT
     50-00-0, Formaldehyde, processes 57-12-5D, Cyanide, compds.
     74-93-1, Methyl mercaptan, processes 75-07-0, Acetaldehyde, processes 75-18-3, Dimethyl sulfide 75-44-5, Phosgene
     75-50-3, Trimethylamine, processes 79-09-4, Propionic acid,
     processes 100-42-5, Styrene, processes 107-92-6, Butyric acid,
     processes 109-52-4, Valeric acid, processes 110-81-6, Diethyl
     disulfide 110-86-1, Pyridine, processes 352-93-2, Diethyl
     sulfide 503-74-2, Iso-valeric acid 505-60-2, Mustard gas 624-92-0 630-08-0, Carbon monoxide, processes 693-07-2, 2-Chloroethyl ethyl sulfide 7440-38-2D, Arsenic, compds.
     7664-41-7, Ammonia, processes
                                        7704-34-9D, Sulfur, compds.
     7783-06-4, Hydrogen sulfide, processes
     RL: ADV (Adverse effect, including toxicity); CPS (Chemical
     process); PEP (Physical, engineering or chemical process); POL
     (Pollutant); REM (Removal or disposal); BIOL (Biological study);
     OCCU (Occurrence); PROC (Process)
        (catalytic compns. for removal of
        contaminants such as warfare agents, and
        materials incorporating these compns.)
IT
     3251-23-8, Copper (II) nitrate 7439-89-6D, Iron,
     complexes with heteropolytungstates 7440-22-4D, Silver, complexes with heteropolytungstates 7440-45-1D, Cerium,
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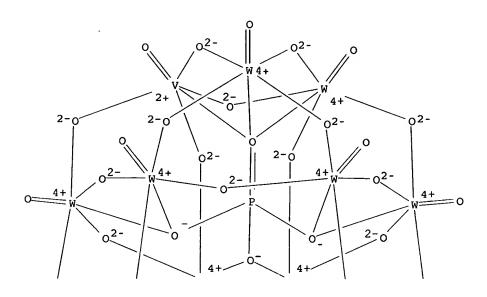
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complexes with heteropolytungstates 10108-73-3, Cerium
     (III) nitrate 10141-05-6, Cobalt (II) nitrate
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     13138-45-9, Nickel (II) nitrate 13770-18-8,
     Copper (II) perchlorate 34946-82-2, Copper (II)
     trifluoromethanesulfonate 38465-60-0, Copper (II)
     tetrafluoroborate
     RL: CAT (Catalyst use); USES (Uses)
        (catalytic compns. for removal of
        contaminants such as warfare agents, and
        materials incorporating these compns.)
L114 ANSWER 13 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
                          2004:769381 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                          142:227445
TITLE:
                          Tuning the formal potentials of new
                          VIV-substituted Dawson-type
                          polyoxometalates for facile synthesis
                          of metal nanoparticles
AUTHOR (S):
                          Keita, Bineta; Mbomekalle, Israel-Martyr;
                          Nadjo, Louis; Haut, Christian
CORPORATE SOURCE:
                          CNRS. Electrochimie et Photoelectrochimie.
                          Laboratoire de Chimie Physique, UMR 8000,
                          Universite Paris-Sud, Orsay, 91405, Fr.
SOURCE:
                          Electrochemistry Communications (2004), 6(10),
                          978-983
                          CODEN: ECCMF9; ISSN: 1388-2481
PUBLISHER:
                          Elsevier B.V.
DOCUMENT TYPE:
                          Journal
LANGUAGE:
                          English
     Dawson-type V-substituted POMs were synthesized and characterized
     both in their oxidized VV and reduced VIV states. They were
     stable in the presence of dioxygen. The apparent formal
     potentials of the V-centers within these mols. can be manipulated
     to span an appreciable potential range through variation of the
     overall atomic composition of the POMs and/or localization of the substituent V-atom in the framework. For each mol., this
     formal potential is pH-independent for pH values larger than 2.5.
     The stability of both forms of these mols. over a large pH domain
     was exploited for the synthesis of metal nanoparticles,
     because this class of POMs presents several interesting features
     not available usually with other chemical reductants. The preparation of
     PdO nanoparticles from [PdCl4]2- was selected as an illustrative
     example. UV-Vis spectroscopy and TEM micrograph were used to
     confirm this successful synthesis.
     12412-90-7P 85585-35-9P 161338-89-2P
     202462-99-5P 258869-02-2P 841244-79-9P
     841244-91-5P 841244-95-9P
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PNU (Preparation, unclassified); PRP (Properties); RCT
     (Reactant); PREP (Preparation); PROC (Process); RACT (Reactant or
     reagent)
        (tuning formal potentials of new VIV-substituted Dawson-type
        polyoxometalates for facile synthesis of metal
        nanoparticles)
    12412-90-7 HCAPLUS
    Vanadate(8-), [dotriaconta-\u03c4-oxoheptadecaoxo[\u03c49-[phosphato(3-
     )-κ0:κ0:κ0:κ0':κ0':κ0'':.kappa
     .0'':κ0''':κ0''']]heptadecatungstate]tetra-μ-
     oxooxo[μ9-[phosphato(3-)-κ0:κ0:κ0:κ0':.
     kappa.0':\kappa0'':\kappa0'':\kappa0''':\kappa0''']]- (9CI)
     (CA INDEX NAME)
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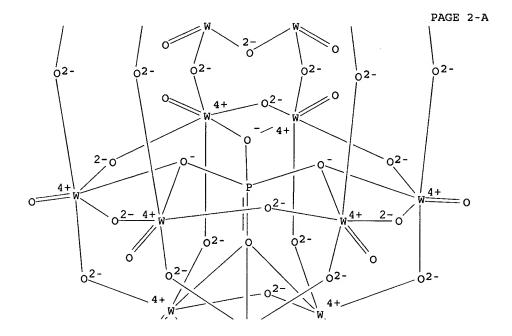
IT

RN

CN

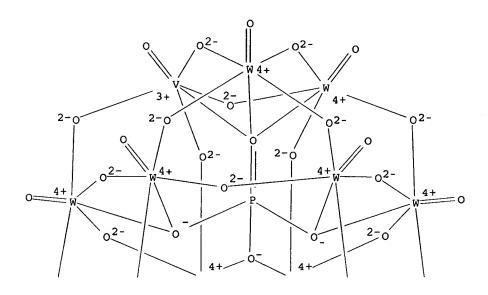
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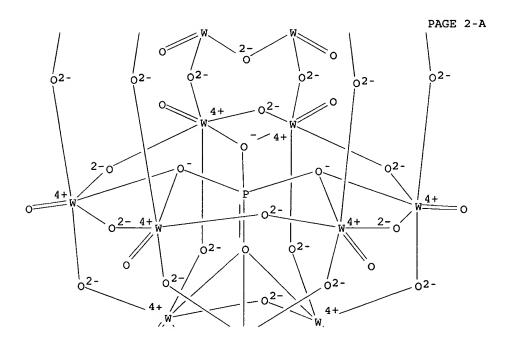




RN 85585-35-9 HCAPLUS
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kappa.0':κ0'':κ0'':κ0''':κ0''']]- (9CI)
(CA INDEX NAME)

PAGE 1-A

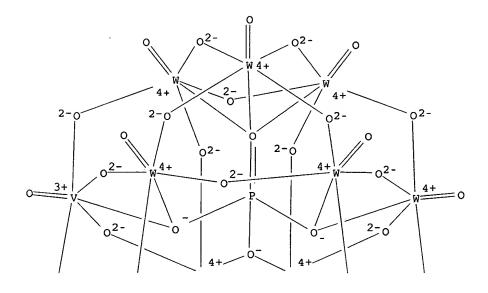


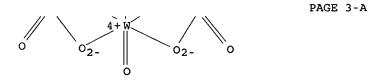


RN 161338-89-2 HCAPLUS
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.0'': κ 0''': κ 0''']]heptadecatungstate]tetra- μ -oxooxo[μ 9-[phosphato(3-)- κ 0: κ 0: κ 0: κ 0':.kappa.0': κ 0'': κ 0'': κ 0''': κ 0''': κ 0''']]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 3-A

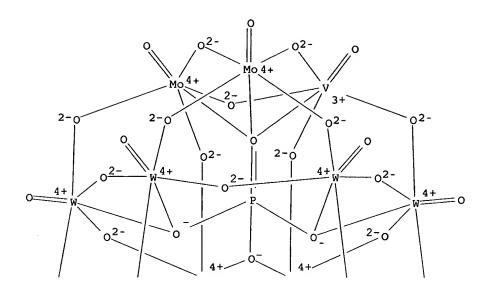


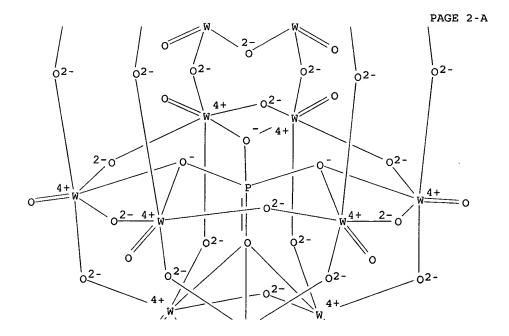


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Les Henderson Page 59 571-272-2538

PAGE 1-A

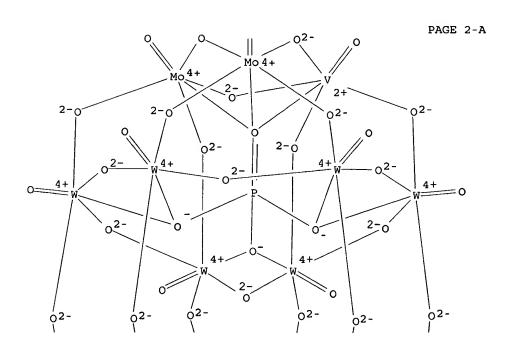




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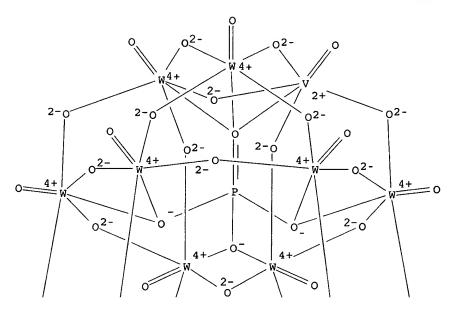
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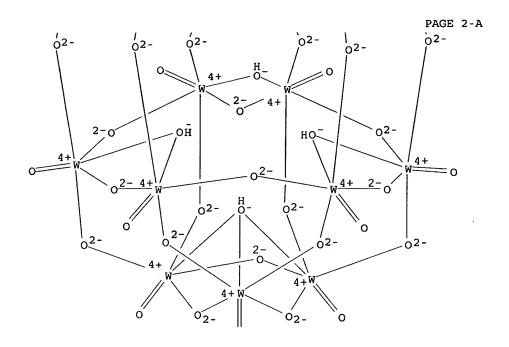
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RN 841244-79-9 HCAPLUS
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'':κΟ''':κΟ''']](tri-μ-hydroxy-μ3hydroxydotriaconta-μ-οχοheptadecaoxoheptadecatungstate)- (9CI)
(CA INDEX NAME)

PAGE 1-A



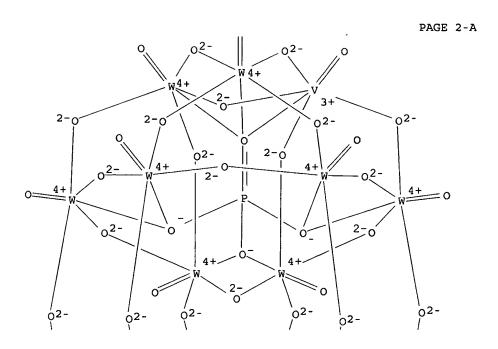


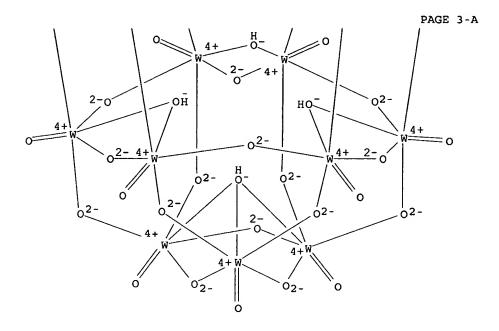
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RN 841244-91-5 HCAPLUS
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'':κΟ'':κΟ'']](tri-μ-hydroxy-μ3hydroxydotriaconta-μ-οχοheptadecaoxoheptadecatungstate)- (9CI)
(CA INDEX NAME)

PAGE 1-A

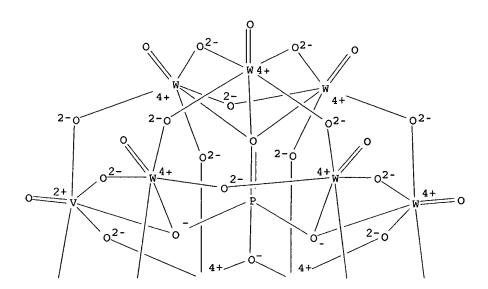
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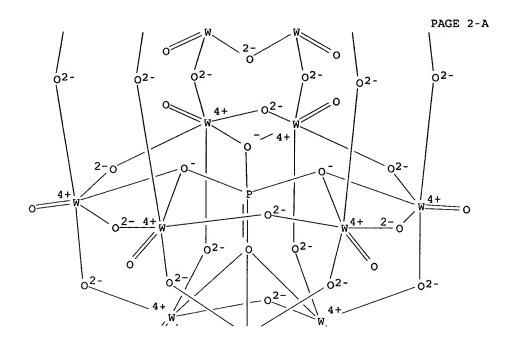


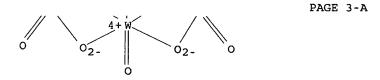


RN 841244-95-9 HCAPLUS
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kappa.0':κ0'':κ0'':κ0''':κ0''']]- (9CI)
(CA INDEX NAME)

PAGE 1-A







CC 72-2 (Electrochemistry)

Section cross-reference(s): 73, 78

ST Dowson type **polyoxometalate** vanadium substituted formal potential nanoparticle palladium

IT Heteropoly acids

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PNU (Preparation, unclassified); PRP (Properties); RCT (Reactant); PREP (Preparation); PROC (Process); RACT (Reactant or reagent)

(Dawson-type; tuning formal potentials of new VIV-substituted Dawson-type polyoxometalates for facile synthesis of metal nanoparticles)

IT Surface structure

(TEM imaging; of palladium nanoparticles formed on vanadium-substituted Dawson-type polyoxometalates with glassy carbon electrode in neutral solns.)

IT Redox reaction

(electrochem.; of new VIV-substituted Dawson-type polyoxometalates)

IT Oxidation potential

UV and visible spectra

(of vanadium-substituted Dawson-type polyoxometalates on glassy carbon electrode in neutral solns.)

IT Cyclic voltammetry

(of vanadium-substituted Dawson-type polyoxometalates with glassy carbon electrode in neutral solns.)

IT Formal potential Nanoparticles

```
(tuning formal potentials of new VIV-substituted Dawson-type
        polyoxometalates for facile synthesis of metal
        nanoparticles)
     7440-44-0, Glassy carbon, uses
TТ
     RL: DEV (Device component use); USES (Uses)
        (glassy; cyclic voltammetry of vanadium-substituted Dawson-type
        polyoxometalates with glassy carbon electrode in
        neutral solns.)
ΙT
     7440-05-3P, Palladium, processes
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PNU (Preparation, unclassified); PREP (Preparation);
     PROC (Process)
        (tuning formal potentials of new VIV-substituted Dawson-type
        polyoxometalates for facile synthesis of metal
        nanoparticles)
     12412-90-7P 85585-35-9P
                              139902-56-0P
TΤ
     161338-89-2P 202462-99-5P 258869-02-2P
     841244-63-1P
                    841244-66-4P
                                    841244-72-2P 841244-79-9P
     841244-82-4P
                    841244-85-7P
                                    841244-88-0P 841244-91-5P
     841244-95-9P
                    841245-02-1P
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PNU (Preparation, unclassified); PRP (Properties); RCT
     (Reactant); PREP (Preparation); PROC (Process); RACT (Reactant or
     reagent)
        (tuning formal potentials of new VIV-substituted Dawson-type
        polyoxometalates for facile synthesis of metal
        nanoparticles)
REFERENCE COUNT:
                                THERE ARE 29 CITED REFERENCES AVAILABLE
                                FOR THIS RECORD. ALL CITATIONS AVAILABLE
                                IN THE RE FORMAT
L114 ANSWER 14 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2004:690229 HCAPLUS
DOCUMENT NUMBER:
                          141:357912
TITLE:
                         Preparation of surface modifications of
                         mesoporous titania with monosubstituted Keggin
                         units and their catalytic performance for
                         organochlorine pesticide and dyes under UV
                          irradiation
AUTHOR(S):
                         Yang, Yu; Guo, Yihang; Hu, Changwen; Wang,
                         Yuanhong; Wang, Enbo
CORPORATE SOURCE:
                          Institute of Polyoxometalate Chemistry,
                         Faculty of Chemistry, Northeast Normal
                         University, Changchun, 130024, Peop. Rep.
                         China
SOURCE:
                         Applied Catalysis, A: General (2004),
                         273 (1-2), 201-210
                         CODEN: ACAGE4; ISSN: 0926-860X
PUBLISHER:
                         Elsevier B.V.
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     A kind of novel and efficient catalyst, mesoporous TiO2 (anatase)
     modified by two transition metal-monosubstituted
     polyoxometalates (POMs), i.e., K5[Ni(H2O)PW11O39] (PW11Ni)
     and K5[Co(H2O)PW11039] (PW11Co), was used to photodegrade an
     organochlorine pesticide, hexachlorobenzene (HCB), and three kinds
     of different dyes under UV irradiation These dyes have various chemical
     structures, either azoic (Congo red (CR), Methyl orange (MO)), or anthraquinonic (Alizarin S (AS)) or fluorescent (Neutral red
     (NR)). Anatase TiO2 was prepared by combined sol-gel and programmed
     temperature hydrothermal methods at a lower temperature (200° C), and
     these as-synthesized TiO2 particles were further functionalized by
     3-aminopropyltriethoxysilane (APS). Amine-functionalized TiO2
     materials impregnated with monosubstituted
     polyoxometalates were prepared by coordination of Ni or Co
     in the units of polyoxometalates with surface amine
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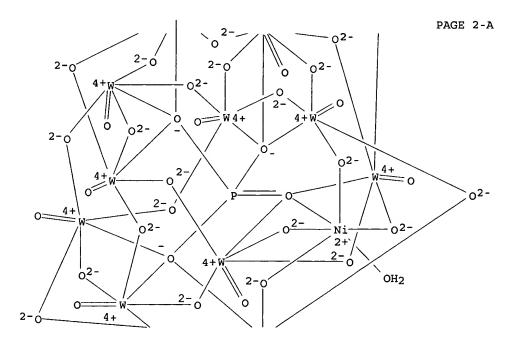
groups in TiO2. The resulting materials have been characterized by several methods, including UV diffuse reflectance spectroscopy (UV/DRS), x-ray diffraction (XRD), inductively coupled plasma atomic emission spectrometry (ICP-AES), 31P magic-angle spinning NMR (MAS NMR), transmission electron microscopy (TEM), and nitrogen adsorption. The conversions of organochlorine pesticide (HCB) and dyes (CR, MO, AS and NR) remarkably increased on UV-irradiating these as-prepared catalysts compared with the results over traditional anatase TiO2; in particular, HCB conversion reached above 98% after UV-irradiating the catalysts for 60 min. With good photocatalytic activity under UV irradiation and the ability to be readily separated from the reaction system, this novel kind of catalyst exhibited the potential to be effective in the treatment of organic pollutants in aqueous systems. 37194-75-5DP, surface reaction product with amine-functionalized titania 39293-41-9DP, surface reaction product with amine-functionalized titania RL: CAT (Catalyst use); PRP (Properties); SPN (Synthetic preparation); PREP (Preparation); USES (Uses) (sol-gel preparation and properties of mesoporous TiO2 modified by transition metal-substituted polyoxometalates and its activity as photocatalyst for degradation of aqueous hexachlorobenzene and dyes) 37194-75-5 HCAPLUS Tungstate(5-), (aquanickelate)tetracosa- μ -oxoundecaoxo[μ 12-[phosphato(3-)- κ 0: κ 0: κ 0: κ 0: κ 0': κ 8 pa.0'κ0'':κ0'':κ0'':κ0''':.kap pa.O''']]undeca-, pentapotassium (9CI) (CA INDEX NAME)

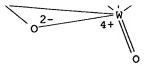
TТ

RN

CN

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





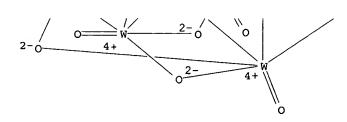
●5 K⁴

RN 39293-41-9 HCAPLUS
CN Tungstate(5-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[phosphato(3-)-κ0:κ0:κ0':κ0':kap
pa.0':κ0'':κ0'':κ0'':κ0'':ka
ppa.0'']]undeca-, pentapotassium (9CI) (CA INDEX NAME)

PAGE 1-A

 \mathbb{N}

PAGE 2-A ი2-<u>02-</u> H20 o 2-0 2-() () 2-0 02 02-2+ 0 2 0= o2-2-0 02-<u>4+</u> 0 02-2-4+ 4+ 02-2-2-Ó 0



PAGE 3-A

●5 K+

CC 74-1 (Radiation Chemistry, Photochemistry, and Photographic and
Other Reprographic Processes)
Section cross-reference(s): 60, 67

ST polyoxometalate surface modification mesoporous titania photocatalyst; organochlorine pesticide water pollutant dye photodegrdn catalyst

IT Reflection spectra

(UV-visible diffuse; sol-gel preparation and properties of mesoporous TiO2 modified by transition metal-substituted polyoxometalates and its activity as photocatalyst for degradation of aqueous hexachlorobenzene and dyes)

IT UV and visible spectra

(diffuse reflection; sol-gel preparation and properties of mesoporous TiO2 modified by transition metal-substituted polyoxometalates and its activity as photocatalyst for degradation of aqueous hexachlorobenzene and dyes)

IT Porous materials

(films, mesoporous; sol-gel preparation and properties of mesoporous TiO2 modified by transition metal-substituted polyoxometalates and its activity as photocatalyst for degradation of aqueous hexachlorobenzene and dyes)

```
TT
     Wastewater treatment
         (photocatalytic; photocatalyst activity of sol-gel derived
        mesoporous TiO2 modified by transition metal-substituted
        polyoxometalates for degradation of aqueous organochlorine
        pesticide and dyes in relation to)
     Catalysis
IT
        (photochem.; sol-gel preparation and properties of mesoporous TiO2
        modified by transition metal-substituted
        polyoxometalates and its activity as photocatalyst for
        degradation of aqueous hexachlorobenzene and dyes)
ΙT
        (porous, mesoporous; sol-gel preparation and properties of
        mesoporous TiO2 modified by transition metal-substituted
        polyoxometalates and its activity as photocatalyst for
        degradation of aqueous hexachlorobenzene and dyes)
TΤ
     Photolysis catalysts
     Pore size distribution
     Sol-gel processing
     X-ray diffraction
        (sol-gel preparation and properties of mesoporous TiO2 modified by
        transition metal-substituted polyoxometalates and its
        activity as photocatalyst for degradation of aqueous hexachlorobenzene
        and dyes)
TΤ
     Heteropoly acids
     RL: CAT (Catalyst use); PRP (Properties); SPN (Synthetic
     preparation); PREP (Preparation); USES (Uses)
        (transition metal tungstophosphates; sol-gel preparation and
        properties of mesoporous TiO2 modified by transition
        metal-substituted polyoxometalates and its activity
        as photocatalyst for degradation of aqueous hexachlorobenzene and dyes)
     13463-67-7DP, Titania, amino-functionalized, surface product with
     transition metal-substituted polyoxometalates
RL: CAT (Catalyst use); PRP (Properties); SPN (Synthetic
     preparation); PREP (Preparation); USES (Uses)
        (anatase-type; sol-gel preparation and properties of mesoporous TiO2
        modified by transition metal-substituted
        polyoxometalates and its activity as photocatalyst for
        degradation of aqueous hexachlorobenzene and dyes)
     118-74-1, Hexachlorobenzene 130-22-3, Alizarin S
TT
     Methyl orange 553-24-2, Neutral red 573-58-0, Congo red RL: RCT (Reactant); RACT (Reactant or reagent)
        (model reaction; photocatalyst activity of sol-gel derived
        mesoporous TiO2 modified by transition metal-substituted polyoxometalates for degradation of aqueous organochlorine
        pesticide and dyes)
TT
     919-30-2DP, 3-Aminopropyltriethoxysilane, surface product with
     titania and polyoxometalates 37194-75-5DP,
     surface reaction product with amine-functionalized titania
     39293-41-9DP, surface reaction product with
     amine-functionalized titania
     RL: CAT (Catalyst use); PRP (Properties); SPN (Synthetic
     preparation); PREP (Preparation); USES (Uses)
        (sol-gel preparation and properties of mesoporous TiO2 modified by
        transition metal-substituted polyoxometalates and its
        activity as photocatalyst for degradation of aqueous hexachlorobenzene
        and dyes)
TΤ
     546-68-9, Titanium tetraisopropoxide
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (sol-gel preparation and properties of mesoporous TiO2 modified by
        transition metal-substituted polyoxometalates and its
        activity as photocatalyst for degradation of aqueous hexachlorobenzene
        and dyes)
REFERENCE COUNT:
                                THERE ARE 44 CITED REFERENCES AVAILABLE
                                FOR THIS RECORD. ALL CITATIONS AVAILABLE
                                IN THE RE FORMAT
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L114 ANSWER 15 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                          2004:453687 HCAPLUS
DOCUMENT NUMBER:
                          141:162820
                          Polyoxometalates on Cationic Silica
TITLE:
                          Nanoparticles. Physicochemical Properties of
                          an Electrostatically Bound Multi-Iron Catalyst
AUTHOR (S):
                          Okun, Nelya M.; Ritorto, Michelle
                          D.; Anderson, Travis M.; Apkarian, Robert P.;
                          Hill, Craig L.
                          Department of Chemistry, Emory University,
CORPORATE SOURCE:
                          Atlanta, GA, 30322, USA
SOURCE:
                          Chemistry of Materials (2004), 16(13),
                          2551-2558
                          CODEN: CMATEX; ISSN: 0897-4756
PUBLISHER:
                          American Chemical Society
DOCUMENT TYPE:
                          Journal
LANGUAGE:
                          English
     Reaction of a solution of the multi-iron polyoxometalate
     (POM) K9[(FeIII(OH2)2)3(A-\alpha-PW9O34)2] (K91) with a colloidal
     suspension of cationic silica nanoparticles ((Si/AlO2)Cl) results
     in the production of a new heterogeneous oxidation catalyst
     (K81/(Si/AlO2)). Dynamic light scattering data, coupled with
     elemental anal. and streaming potential measurements suggests that
     there are 58 mols. of POM electrostatically bound to the surface
     of each silica particle on average Transmission electron microscopy
     confirms the presence of the POM on the surface of the cationic silica and shows the diameter of the (Si/AlO2)Cl and of the
     K81/(Si/AlO2) nanoparticles to be .apprx.12 and .apprx.17 nm,
            Significantly, cryo-high-resolution SEM (cryo-HRSEM) shows
     that the POM retards the natural gelation process that colloidal
     silica is known to undergo upon aging. EPR and catalytic data collectively suggest that the exchange of the cationic silica for
     one of the nine K+ cations associated with each POM is responsible
     for subtle structural changes in the POM which result in its
     activation as a catalyst.
CC
     67-1 (Catalysis, Reaction Kinetics, and Inorganic Reaction
     Mechanisms)
     Section cross-reference(s): 66
     polyoxometalate cationic silica nanoparticle property
     electrostatically bound multiiron catalyst
TΤ
     Colloids
     Nanoparticles
     Oxidation catalysts
     Surface structure
        (polyoxometalates on cationic silica nanoparticles
        and physicochem. properties of electrostatically bound
        multi-iron catalyst)
TТ
     Heteropoly acids
     RL: CAT (Catalyst use); PRP (Properties); USES (Uses)
        (polyoxometalates on cationic silica nanoparticles
        and physicochem. properties of electrostatically bound
        multi-iron catalyst)
TT
     7631-86-9, Silica, uses
                                728945-74-2
     RL: CAT (Catalyst use); PRP (Properties); USES (Uses)
        (polyoxometalates on cationic silica nanoparticles
        and physicochem. properties of electrostatically bound
        multi-iron catalyst)
REFERENCE COUNT:
                                THERE ARE 60 CITED REFERENCES AVAILABLE
                          60
                                FOR THIS RECORD. ALL CITATIONS AVAILABLE
                                IN THE RE FORMAT
L114 ANSWER 16 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                          2004:304574 HCAPLUS
DOCUMENT NUMBER:
                          141:307138
TITLE:
                          Study of some polyoxometallates of
```

Keggin's type as potential antitumor agents

Holclajtner-Antunovic, Ivanka; Kuntic, Vesna; AUTHOR(S): Juranic, Zorica; Filipovic, Ivana; Mioc, Ubavka; Stanojkovic, Tatjana; Zizak, Zeljko University School of Physical Chemistry, CORPORATE SOURCE: Belgrade, Yugoslavia SOURCE: Jugoslovenska Medicinska Biohemija (2004), 23(1), 25-30 CODEN: JMBIFF; ISSN: 0354-3447 PUBLISHER: Drustvo Medicinskih Biohemicara Jugoslavije Journal DOCUMENT TYPE: LANGUAGE: English/Croatian The antitumor action of three polyoxometallate compds. of Keggin's type: 12-molibdophosphoric acid (MoPA), 12-tungstophosphoric acid (WPA) and Mg salt of WPA (MgHWPA) was studied in vitro. For human cervix carcinoma (HeLa) cells survival, as well as for nonstimulated and stimulated peripheral blood mononuclear cells (PBMC), MTT test was applied and IC50 values of POMs were determinated. Index selectivity for WPA and MgHWPA are 1.9 and 1.8, calculated for nonstimulated, as well as 2.5 and 2.0, calculated for stimulated PBMS. Combination of studied POMs do not contribute to their lower IC50 values. Apoptosis detection implies mild cytotoxic effect of WPA and more cytostatical effect of MgHWPA. Combination of each of the studied POMs with caffeine decreases HeLa survival in dose dependent way. None of the studied POMs in the used concns. (up to 100 µmol/L) damages blood cells and/or decreases their number TТ 105814-03-7 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (POM of keggin's type, MgHWPA acted cytostatically on stimulated and non-stimulated PBMC, HeLa cell and

combination with caffeine decreased survival in human

Tungstate (3-), tetracosa- μ -oxododecaoxo [μ 12-[phosphato(3-)-

a-, magnesium hydrogen (1:1:1) (9CI) (CA INDEX NAME)

cervix carcinoma cells)

':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec

105814-03-7 HCAPLUS

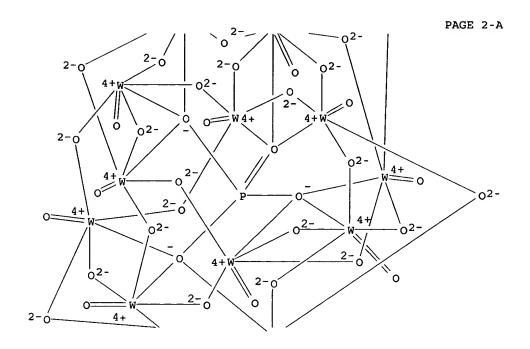
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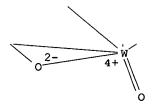
RN

CN

PAGE 1-A

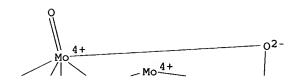
0 W4+ W4+ 02-

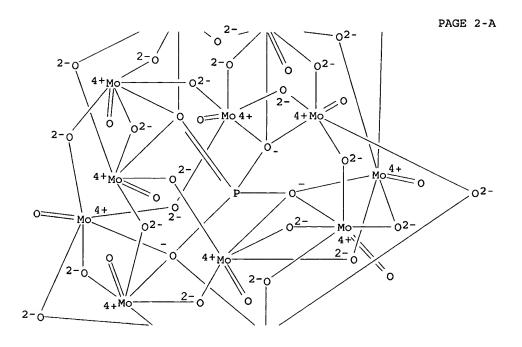


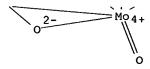


● H+

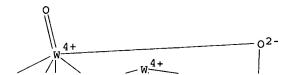
● Mg²⁺

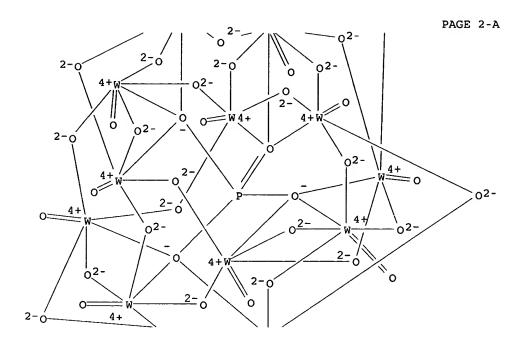


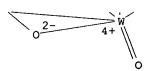




●3 H+







●3 H+

CC 1-6 (Pharmacology)

ST polyoxometallate keggin molibdophosphoric

tungstophosphoric acid antitumor apoptosis cervix carcinoma

IT Antitumor agents

Human

(POMs of keggin's type showed no considerable effect on survival with WPA showing mild cytotoxic effect, MgHWPA being more cytostatic, MoPA with insignificant effect and their combination with caffeine decreasing survival in human HeLa cell)

IT Uterus, neoplasm

(cervix, carcinoma; POMs of keggin's type showed no considerable effect on survival with WPA showing mild cytotoxic effect, MgHWPA being more cytostatic, MoPA with insignificant effect and their combination with caffeine decreasing survival in human HeLa cell)

IT Carcinoma

(cervix; POMs of keggin's type showed no considerable effect on survival with WPA showing mild cytotoxic effect, MgHWPA being more cytostatic, MoPA with insignificant effect and their combination with caffeine decreasing survival in human HeLa cell)

IT 105814-03-7

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL

(Biological study); USES (Uses) (POM of keggin's type, MgHWPA acted cytostatically on stimulated and non-stimulated PBMC, HeLa cell and combination with caffeine decreased survival in human cervix carcinoma cells) ΙT 12026-57-2, 12-Molybdophosphoric acid RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (POM of keggin's type, MoPA showed insignificant antiproliferative effect on stimulated and non-stimulated PBMC, HeLa cell and combination with caffeine decreased survival in human cervix carcinoma cells) 1343-93-7, 12-Tungstophosphoric acid RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (POM of keggin's type, WPA showed mild antiproliferative effect on stimulated and non-stimulated PBMC, HeLa cell and combination with caffeine decreased survival in human cervix carcinoma cells) REFERENCE COUNT: 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L114 ANSWER 17 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2004:271023 HCAPLUS DOCUMENT NUMBER: 141:56014 TITLE: Effects of support on bifunctional methanol oxidation pathways catalyzed by polyoxometallate Keggin clusters Liu, Haichao; Iglesia, Enrique E.O. Lawrence Berkeley National Laboratory, AUTHOR(S): CORPORATE SOURCE: Chemical Sciences Division, Department of Chemical Engineering, University of California at Berkeley, Berkeley, CA, 94720, USA Journal of Catalysis (2004), 223(1), 161-169 CODEN: JCTLA5; ISSN: 0021-9517 SOURCE: PUBLISHER: Elsevier Science DOCUMENT TYPE: Journal LANGUAGE: English OTHER SOURCE(S): CASREACT 141:56014 H5PV2Mo10040 polyoxometallate Keggin clusters supported oxidation reactions to form HCHO, Me formate (MF), and dimethoxymethane (DMM). The rate and selectivity and the of the oxide supports. Raman spectroscopy showed that Keggin structures remained essentially intact on ZrO2, TiO2, and SiO2 after treatment in air at 553 K, but decomposed to MoOx and VOx

on ZrO2, TiO2, SiO2, and Al2O3 are effective catalysts for CH3OH structure of supported clusters depend on the surface properties oligomers on Al2O3. Accessible protons per Keggin unit (KU) were measured during CH3OH oxidation by titration with 2,6-di-tert-Bu pyridine. For similar KU surface d. (0.28 - 0.37 KU/nm2), the number of accessible protons was larger on SiO2 than on ZrO2 and TiO2 and much smaller on Al2O3 supports, even though residual di-Me ether (DME) synthesis rates after titrant saturation indicated that the fractional dispersion of KU was similar on the first three supports. These effects of support on structure and on H+ accessibility reflect varying extents of interaction between polyoxometallate clusters and supports. Rates of CH3OH oxidative dehydrogenation per KU were higher on ZrO2 and TiO2 than on SiO2 at similar KU surface d. (0.28 - 0.37 KU/nm2) and dispersion, indicating that redox properties of Keggin clusters depend on the identity of the support used to disperse them. ZrO2 and TiO2 supports appear to enhance the reducibility of anchored polyoxometallate clusters. Rates were much lower on Al203, because structural degradation led to less reactive MoOx and VOx domains. CH3OH reactions involve primary oxidation to

form HCHO and subsequent secondary reactions to form DMM and MF. These reactions involve HCHO-CH3OH acetalization steps leading to methoxymethanol (CH3OCH2OH) or hemiacetal intermediates, which condense with CH3OH on acid sites to form DMM or dehydrogenate to form MF. The COx formation rate is much lower than that of other reactions, and DME forms in parallel pathways catalyzed by acid sites. Secondary reactions leading to DMM and MF are strongly influenced by the support surface. Acidic SiO2 surfaces favored DMM formation, while amphoteric or dehydrogenating surfaces on ZrO2 and TiO2 led to MF formation, as a result of the varying role of each support in directing the reactions of HCHO and CH3OH and of the CH3OCH2OH intermediates toward DMM or MF, which was confirmed using phys. catalyst-pure support mixts. The reaction pathways are consistent with the effects of residence time and of the partial removal of H+ sites by titration using 2,6-di-tert-Bu pyridine. 12293-21-9, Molybdovanadophosphoric acid (H5Mo10V2PO40) RL: CAT (Catalyst use); USES (Uses) (oxidation catalyst; effects of oxide support acidity and surface structure on selectivity of Mo-V-P-O Keggin cluster catalyst in methanol oxidation rate and mechanism) 12293-21-9 HCAPLUS $\label{eq:Vanadate} \textit{Vanadate(5-), (heptadeca-μ-oxodecaoxodecamolybdate) hepta-μ-}$ oxodioxo [μ12- (phosphato (3-)-κ0:κ0:κ0:κ0 1:K01:K01:K011:K011:K011:K0111

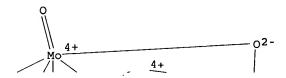
 $\kappa(0''':\kappa(0''')]$ di-, pentahydrogen (9CI) (CA INDEX

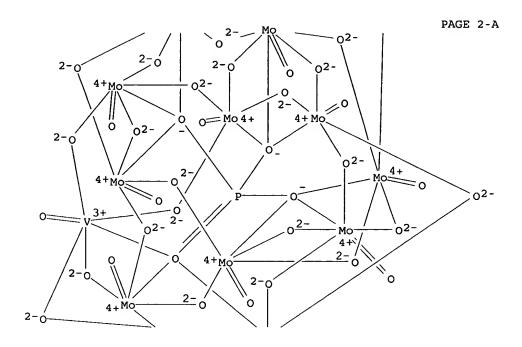
IT

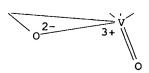
RN

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NAME)







●5 H+

CC 45-4 (Industrial Organic Chemicals, Leather, Fats, and Waxes)

Section cross-reference(s): 23, 67

IT 12293-21-9, Molybdovanadophosphoric acid (H5Mo10V2PO40)

RL: CAT (Catalyst use); USES (Uses)

(oxidation catalyst; effects of oxide support acidity and surface structure on selectivity of Mo-V-P-O Keggin cluster catalyst in methanol oxidation rate and mechanism)

REFERENCE COUNT:

THERE ARE 25 CITED REFERENCES AVAILABLE 25 FOR THIS RECORD. ALL CITATIONS AVAILABLE

IN THE RE FORMAT

L114 ANSWER 18 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2004:229264 HCAPLUS

TITLE: Multifunctional nanomaterials for catalytic decontamination and detection

AUTHOR(S): Hill, Craig L.; Neiwert, Wade A.; Okun, Nelya M.; Anderson, Travis M.; Ritorto, Michelle D.; Han, Jong Woo

CORPORATE SOURCE: Department of Chemistry, Emory University,

Atlanta, Atlanta, GA, 30322, USA

SOURCE:

Abstracts of Papers, 227th ACS National Meeting, Anaheim, CA, United States, March 28-April 1, 2004 (2004), POLY-629. American

Chemical Society: Washington, D. C.

CODEN: 69FGKM

DOCUMENT TYPE: Conference; Meeting Abstract

LANGUAGE: English

New types of materials based on transition-metal oxygen-anion clusters (polyoxometalates or "POMs") have been developed for the detection and decontamination of a range of toxic mols. from indoor air pollutants to chemical warfare agents. Detection in most cases involves an oxidation of the toxic mol. and reduction of the POM with an attendant color change; catalytic decontamination usually involves this step followed by reoxidn. by O2 under ambient conditions. The first prototype material is a nanodomain comprised of 72 bridging catalytically active V6013 units, each connected by 2 tri-triols via triester linkages. material gels toxic agents, detects some by color change and catalyzes their oxidative degradation using air. Three other prototype detecting and/or decontaminating materials are POMs electrostatically bound to cationic fabrics ("catalytic cotton"), esterified-POM-based coordination polymers and POMs bridged by

L114 ANSWER 19 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2004:105152 HCAPLUS 141:234961

DOCUMENT NUMBER:

TITLE: Inhibition of influenza virus infections in

redox-active-metal counterions into microporous structures.

MDCK cells by rare earth borotungstates heteropoly blues with mixed valence

AUTHOR(S): Liu, Jie; Mei, Wen-Jie; Li, An-Xing; Tan,

Cai-Ping; Shi, Shuo; Ji, Liang-Nian

Key Lab. Gene Eng. of Education Ministry,
Coll. Chem. Chem. Eng., Zhongshan Univ., CORPORATE SOURCE:

Guangzhou, 510275, Peop. Rep. China

Gaodeng Xuexiao Huaxue Xuebao (2004), 25(1),

CODEN: KTHPDM; ISSN: 0251-0790

Gaodeng Jiaoyu Chubanshe

DOCUMENT TYPE: Journal LANGUAGE: Chinese

SOURCE:

PUBLISHER:

Transition metal-substituted mixed-valence rare earth borotungstate heteropoly blues, Ln2H3[BWVI9WV2Co(H2O)O39]·n H2O (Ln = La, Ce, Pr, Nd, Sm, Eu or Gd), were prepared and characterized by IR, UV, CV, XPS and electrochem. The cell toxicity and antiviral activity of these rare earth borotungstate heteropoly blues were investigated on the inhibitory effect against influenza (A/H1N1/Jingfang/1/91, A/H3N2/Jingfang/30/95, B/Hufang/1/87) in MDCK cells, and the results suggest that these complexes exhibit an significantly inhibitory activity, which are comparable with those obtained from virazole, against influenza virus infection, and no cytotoxicity is shown on normal cells. The correlation (relation) between the structure of these complexes and their anti-virus activities was also discussed.

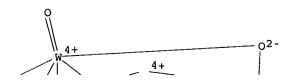
745826-21-5P 745826-22-6P 745826-23-7P 745826-27-1P 745826-30-6P 745826-36-2P 745826-41-9P

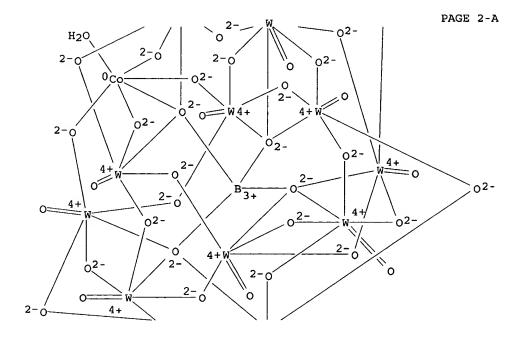
> RL: BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation) (preparation and antiviral activity of cobalt-substituted mixed-valence rare earth borotungstate heteropoly blue complexes)

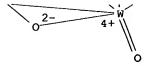
745826-21-5 HCAPLUS ΡN

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PAGE 1-A



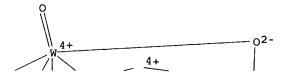


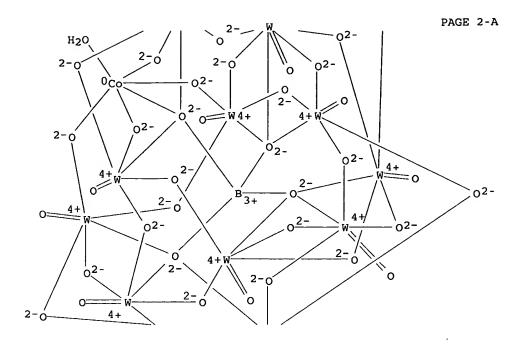


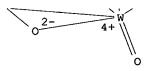
●3 H+

●2 La(III) 3+

RN 745826-22-6 HCAPLUS
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a.O':κΟ':κΟ'':κΟ'':κΟ'':.kappa
.O''':κΟ''']]undeca-, cerium(3+) hydrogen (1:2:3) (9CI) (CA
INDEX NAME)







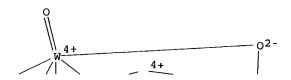
●2 Ce(III) 3+

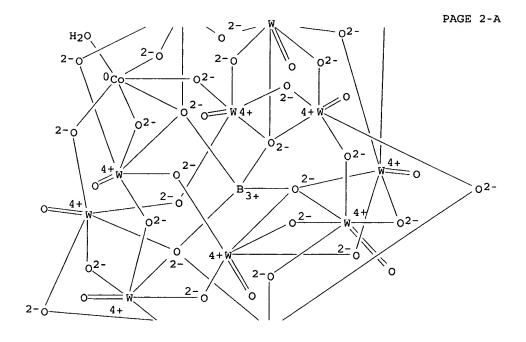
●3 H+

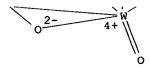
RN 745826-23-7 HCAPLUS

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PAGE 1-A



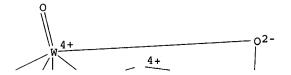


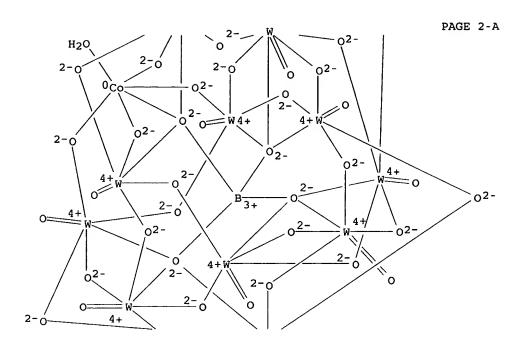


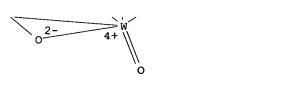
●3 H+

●2 Pr(III) 3+

RN 745826-27-1 HCAPLUS
CN Tungstate(9-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ:κΟ':.kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':.kappa
.O''':κΟ''']]undeca-, neodymium(3+) hydrogen (1:2:3) (9CI)
(CA INDEX NAME)



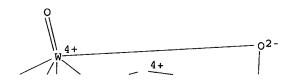


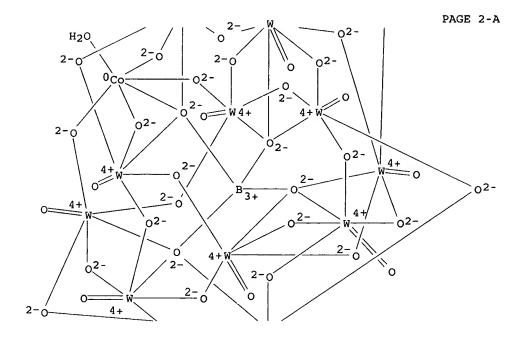


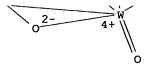
●3 H+

●2 Nd(III) 3+

RN 745826-30-6 HCAPLUS
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a.O':κΟ':κΟ'':κΟ'':κΟ'':.kappa
.O''':κΟ''']]undeca-, samarium(3+) hydrogen (1:2:3) (9CI)
(CA INDEX NAME)





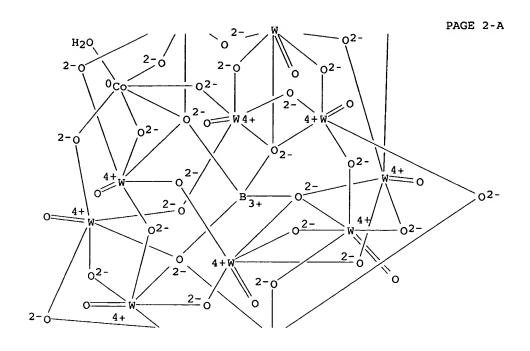


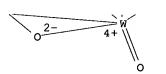
●3 H+

●2 Sm(III) 3+

RN 745826-36-2 HCAPLUS
CN Tungstate(9-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ:κΟ':.kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':.kappa
.O''':κΟ''']]undeca-, europium(3+) hydrogen (1:2:3) (9CI)
(CA INDEX NAME)







●2 Eu(III) 3+

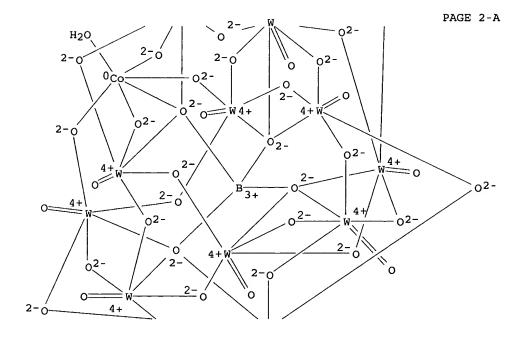
●3 H+

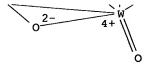
RN

745826-41-9 HCAPLUS
Tungstate(9-), (aquacobaltate)tetracosa- μ -oxoundecaoxo[μ 12-[tetrahydroxyborato(5-)- κ 0: κ 0: κ 0: κ 0': κ CN (CA INDEX NAME)

PAGE 1-A



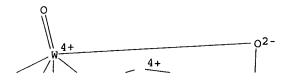


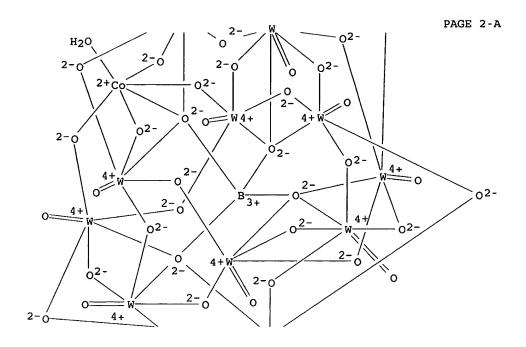


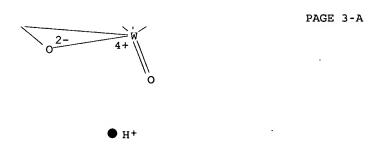
●2 Gd(III) 3+

●3 H+

IT 745826-12-4P 745826-14-6P 745826-16-8P 745826-17-9P 745826-18-0P 745826-19-1P 745826-20-4P RL: BSU (Biological study, unclassified); RCT (Reactant); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent)
(preparation and antiviral activity of cobalt-substituted mixed-valence rare earth borotungstate heteropoly blues complexes) 745826-12-4 HCAPLUS RN CN Tungstate(7-), (aquacobaltate)tetracosa- μ -oxoundecaoxo[μ 12-[tetrahydroxyborato(5-)-κ0:κ0:κ0:κ0':.kapp a.0':κ0':κ0'':κ0'':κ0'':k0'':.kappa .0''':κ0''']]undeca-, lanthanum(3+) hydrogen (1:2:1) (9CI) (CA INDEX NAME)



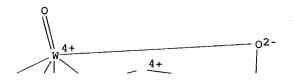


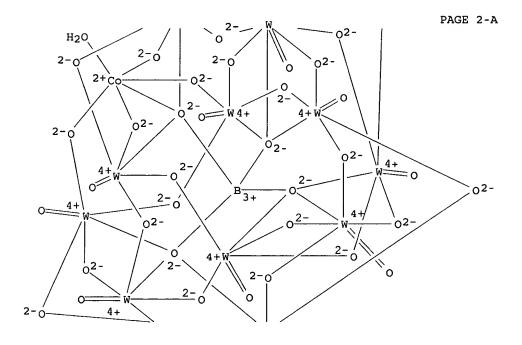


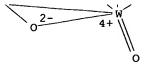
●2 La(III) 3+

RN 745826-14-6 HCAPLUS
CN Tungstate(7-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κ0:κ0:κ0':.kapp
a.0':κ0'':κ0'':κ0'':κ0'':.kappa
.0''':κ0''']]undeca-, cerium(3+) hydrogen (1:2:1) (9CI) (CA INDEX NAME)

PAGE 1-A



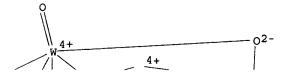


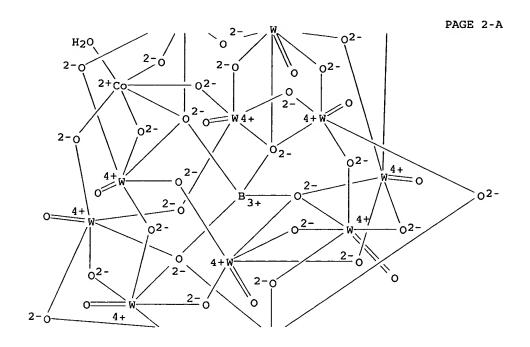


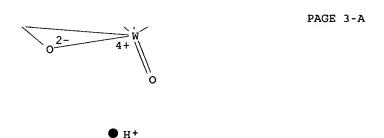
●2 Ce(III) 3+

● H+

RN 745826-16-8 HCAPLUS
CN Tungstate(7-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ:κ:kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':kappa
.O''':κΟ''']]undeca-, praseodymium(3+) hydrogen (1:2:1)
(9CI) (CA INDEX NAME)

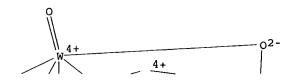


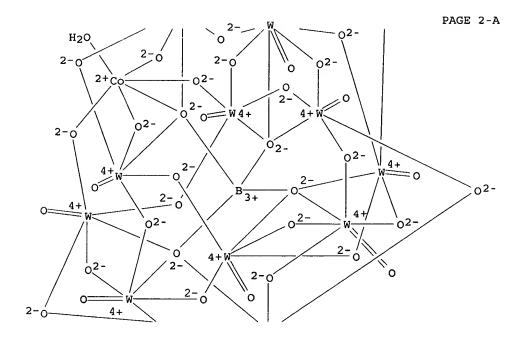


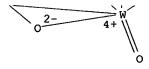


•2 Pr(III) 3+

RN 745826-17-9 HCAPLUS
CN Tungstate(7-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ:κΟ':.kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':κΟ'':.kappa
.O'':κΟ'']]undeca-, neodymium(3+) hydrogen (1:2:1) (9CI)
(CA INDEX NAME)



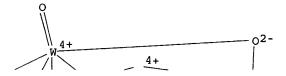


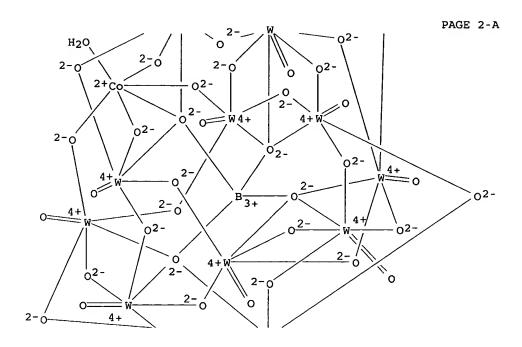


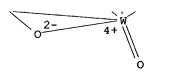
● H+

●2 Nd(III) 3+

RN 745826-18-0 HCAPLUS
CN Tungstate(7-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ:κΟ':.kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':.kappa
.O''':κΟ''']]undeca-, samarium(3+) hydrogen (1:2:1) (9CI)
(CA INDEX NAME)





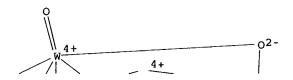


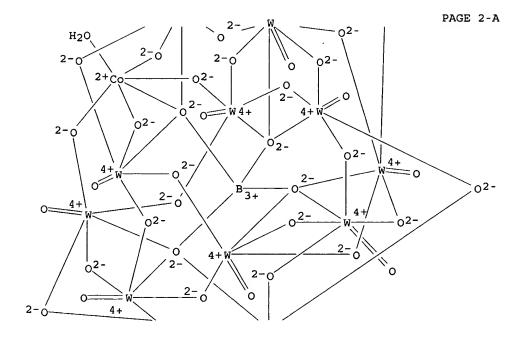
● H+

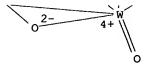
●2 Sm(III) 3+

RN 745826-19-1 HCAPLUS
CN Tungstate(7-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ:κΟ':.kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':.kappa
.O''':κΟ''']]undeca-, europium(3+) hydrogen (1:2:1) (9CI)
(CA INDEX NAME)

PAGE 1-A



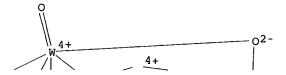


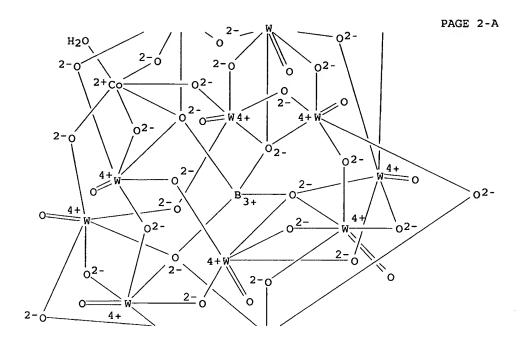


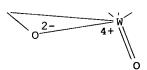
●2 Eu(III) 3+

● H+

RN 745826-20-4 HCAPLUS
CN Tungstate(7-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ:κΟ':.kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':κΟ'':.kappa
.O''':κΟ''']]undeca-, gadolinium(3+) hydrogen (1:2:1) (9CI)
(CA INDEX NAME)







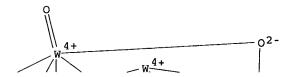
●2 Gd(III) 3+

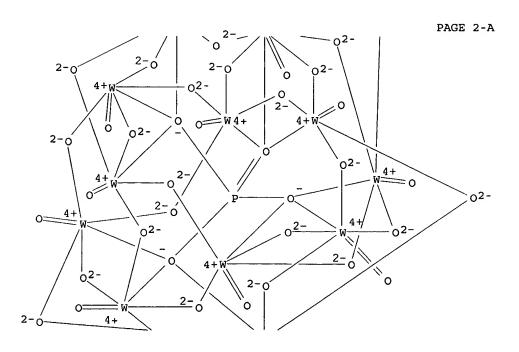
● H+

78-7 (Inorganic Chemicals and Reactions) CC Section cross-reference(s): 1, 4, 10 IT Animal cell line (MDCK; antiviral activity of cobalt-substituted mixed -valence rare earth borotungstate heteropoly blue complexes) IT Antiviral agents Influenza virus (antiviral activity of cobalt-substituted mixed -valence rare earth borotungstate heteropoly blue complexes) ΙT Reduction, electrochemical (preparation of cobalt-substituted mixed-valence rare earth borotungstate heteropoly blue complexes by) IT Rare earth complexes RL: BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation) (tungstoboric acid complexes; antiviral activity of cobalt-substituted mixed-valence rare earth borotungstate heteropoly blue complexes) ΙT Heteropoly acids

RL: BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation) (tungstoboric, rare earth complexes; antiviral activity of cobalt-substituted mixed-valence rare earth borotungstate heteropoly blue complexes) ΙT 745826-21-5P 745826-22-6P 745826-23-7P 745826-27-1P 745826-30-6P 745826-36-2P 745826-41-9P RL: BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation) (preparation and antiviral activity of cobalt-substituted mixed-valence rare earth borotungstate heteropoly blue complexes) TT 745826-12-4P 745826-14-6P 745826-16-8P 745826-17-9P 745826-18-0P 745826-19-1P 745826-20-4P RL: BSU (Biological study, unclassified); RCT (Reactant); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent) (preparation and antiviral activity of cobalt-substituted mixed-valence rare earth borotungstate heteropoly blues complexes) L114 ANSWER 20 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2003:922792 HCAPLUS DOCUMENT NUMBER: 140:116375 TITLE: Photocatalysis by Titanium Dioxide and Polyoxometalate/TiO2 Cocatalysts. Intermediates and Mechanistic Study AUTHOR(S): Chen, Chuncheng; Lei, Pengxiang; Ji, Hongwei; Ma, Wanhong; Zhao, Jincai; Hidaka, Hisao; Serpone, Nick CORPORATE SOURCE: Laboratory of Photochemistry, Center for Molecular Science, Institute of Chemistry, Chinese Academy of Sciences, Beijing, 100080, Peop. Rep. China SOURCE: Environmental Science and Technology (2004), 38(1), 329-337 CODEN: ESTHAG; ISSN: 0013-936X American Chemical Society PUBLISHER: DOCUMENT TYPE: Journal LANGUAGE: English A representative polyoxometalate, α-12tungstophosphatic acid (PW123-, POM), is loaded on the surface of TiO2 particles used as a cocatalyst to gain further insights into the underlying reaction mechanism and to estimate the feasibility of using the new POM/TiO2 cocatalyst in the photocatalytic degradation of 2,4-dichlorophenol (DCP) in aqueous media. Loading the PW123- species on the surface of TiO2 enhances charge separation in the UV-illuminated TiO2, thereby accelerating the hydroxylation of the initial DCP substrate but not the mineralization of DCP, which is somewhat suppressed in the presence of the polyoxometalate. An increase in the load of POM increases the concentration of aromatic intermediates, and more toxic intermediates, such as 2,6-dichlorodibenzo-p-dioxin, 2,4,6-trichlorophenol, are detected in the PW123-/TiO2 system. By contrast, cleavage of the whole conjugated structure of DCP predominates in TiO2-only dispersions. Strong ESR signals for the superoxide radical anionic species, 02-• (HO2• radicals in acidic media; pH <5) are detected in TiO2-only dispersions; signals of O2- • are much weaker in the TiO2/POM composite system under otherwise identical conditions. Exptl. results imply that enhancement of charge separation in TiO2 photocatalysis does not always result in improvement of the efficiency of mineralization of organic substrates, and the

reaction between organic radical cations and the formed superoxide radical anions may be responsible for the mineralization of the

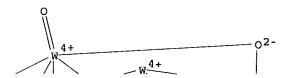


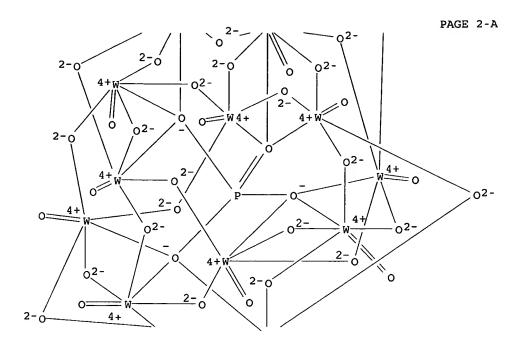


0²⁻4+W

PAGE 3-A

●3 H+







CC 60-2 (Waste Treatment and Disposal)

Section cross-reference(s): 61, 74

ST trichlorophenol photocatalytic degrdn titania

polyoxometalate cocatalyst intermediate mechanism

IT Photolysis

(catalytic; intermediates and mechanism in photocatalysis by titania and polyoxometalate/TiO2 cocatalysts of

2,4-dichlorophenol)

IT Radicals, reactions

RL: FMU (Formation, unclassified); RCT (Reactant); FORM (Formation, nonpreparative); RACT (Reactant or reagent)

(formation and reactions of; intermediates and mechanism in

photocatalysis by titania and polyoxometalate/TiO2 cocatalysts of 2,4-dichlorophenol)

IT Electron transfer

Photolysis catalysts

(intermediates and mechanism in photocatalysis by titania and polyoxometalate/TiO2 cocatalysts of 2,4-dichlorophenol)

IT UV and visible spectra

(of catalysts; intermediates and mechanism in photocatalysis by titania and polyoxometalate/TiO2 cocatalysts of

2,4-dichlorophenol)

IT Wastewater treatment

(oxidation, catalytic, photochem.; intermediates and mechanism in photocatalysis by titania and polyoxometalate/TiO2

```
cocatalysts of 2,4-dichlorophenol in relation to)
IT
     Decomposition
        (photocatalytic; intermediates and mechanism in photocatalysis
        by titania and polyoxometalate/TiO2 cocatalysts of
        2,4-dichlorophenol)
     Wastewater treatment
TΤ
     Water purification
        (photocatalytic; intermediates and mechanism in photocatalysis
        by titania and polyoxometalate/TiO2 cocatalysts of
        2,4-dichlorophenol in relation to)
IT
     Water purification
        (photooxidn., catalytic; intermediates and mechanism in
        photocatalysis by titania and polyoxometalate/TiO2
        cocatalysts of 2,4-dichlorophenol in relation to)
     Oxidation catalysts
TT
        (photooxidn.; intermediates and mechanism in photocatalysis by
        titania and polyoxometalate/TiO2 cocatalysts of
        2,4-dichlorophenol)
ΙT
     88-06-2, 2,4,6-Trichlorophenol 137-19-9, 4,6-Dichloro-1,3-
     benzenediol
                   2138-22-9, 1,2-Benzenediol, 4-chloro-
                                                            3170-83-0,
     Hydroperoxy
                   7722-84-1, Hydrogen peroxide (H2O2), reactions
     11062-77-4, Superoxide 13673-92-2, 3,5-Dichloro-1,2-benzenediol
     RL: FMU (Formation, unclassified); RCT (Reactant); FORM
     (Formation, nonpreparative); RACT (Reactant or reagent)
        (formation and reactions of; intermediates and mechanism in
        photocatalysis by titania and polyoxometalate/TiO2
        cocatalysts of 2,4-dichlorophenol)
TΤ
     33857-26-0, 2,7-Dichlorodibenzodioxin
     RL: POL (Pollutant); OCCU (Occurrence)
        (formation and reactions of; intermediates and mechanism in
        photocatalysis by titania and polyoxometalate/TiO2
        cocatalysts of 2,4-dichlorophenol)
IT
     615-67-8, 2-Chlorohydroquinone
     RL: RCT (Reactant); REM (Removal or disposal); PROC (Process);
     RACT (Reactant or reagent)
        (formation and reactions of; intermediates and mechanism in
        photocatalysis by titania and polyoxometalate/TiO2
        cocatalysts of 2,4-dichlorophenol)
TТ
     1343-93-7 12534-77-9
                           13463-67-7, Titanium
     dioxide, uses
     RL: CAT (Catalyst use); USES (Uses)
        (intermediates and mechanism in photocatalysis by titania and
        polyoxometalate/TiO2 cocatalysts of 2,4-dichlorophenol)
IT
     120-83-2, 2,4-Dichlorophenol
     RL: RCT (Reactant); REM (Removal or disposal); PROC (Process);
     RACT (Reactant or reagent)
        (intermediates and mechanism in photocatalysis by titania and
        polyoxometalate/TiO2 cocatalysts of 2,4-dichlorophenol)
REFERENCE COUNT:
                         62
                               THERE ARE 62 CITED REFERENCES AVAILABLE
                               FOR THIS RECORD. ALL CITATIONS AVAILABLE
                               IN THE RE FORMAT
L114 ANSWER 21 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
                         2003:913046 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         139:392437
TITLE:
                         Materials for degrading contaminants
INVENTOR(S):
                         Okun, Nelya; Hill, Craig L.
PATENT ASSIGNEE(S):
                         Emory University, USA
                         PCT Int. Appl., 34 pp.
SOURCE .
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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Les Henderson Page 107 571-272-2538

APPLICATION NO.

DATE

PATENT NO.

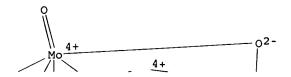
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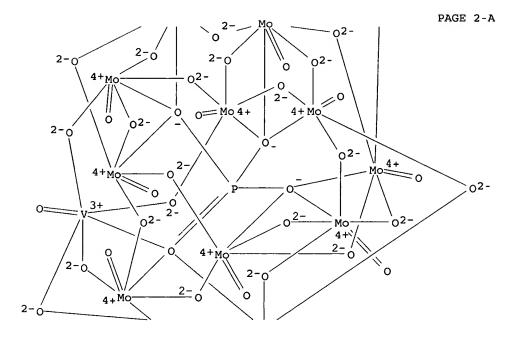
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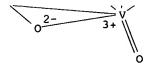
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WO 2003094977
                                      20031120 WO 2003-US14375
                             A2
                                                                               2003
                                                                               0505
                                     20040708
     WO 2003094977
                              A3
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               KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK,
               MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD,
          SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ,
               DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL,
               PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN,
               GQ, GW, ML, MR, NE, SN, TD, TG
     US 2005159307
                              A1
                                      20050721
                                                   US 2003-512336
                                                                               2003
                                                                               0505
PRIORITY APPLN. INFO.:
                                                    US 2002-377740P
                                                                               2002
                                                                               0503
                                                    WO 2003-US14375
                                                                               2003
                                                                               0505
AΒ
     Embodiments of the present invention includes compns.,
     materials including the compns., methods of using the
     compns., and methods of degrading contaminants.
     The composition can include a polyoxometalate
     /cationic silica material. In addition, the compns. can be
     made of a polyoxometalate/cationic silica material, a
     copper (II) salt having a weakly bound anion, and a nitrate salts.
     Further, the compns. can be made of a
     polyoxometalate/cationic silica material, a copper (II)
     salt having a weakly bound anion, a compound selected from
     tetraethylammonium (TEA) nitrate, tetra-n-butylammonium (TBA)
     nitrate, and combinations thereof.
TΤ
     625830-47-9
     RL: NUU (Other use, unclassified); RCT (Reactant); RACT (Reactant
     or reagent); USES (Uses)
         (cationic catalyst support; materials for degrading
         contaminants)
     625830-47-9 HCAPLUS
RN
     Vanadate (5-), (heptadeca-\mu-oxodecaoxodecamolybdate) hepta-\mu-
CN
     oxodioxo [\mu12-[phosphato(3-)-\kappa0:\kappa0:\kappa0:\kappa0
      ':κ0':κ0':κ0'':κ0'':κ0'':κ0''
      :κO''':κO''']]di-, tetrasodium (9CI) (CA INDEX NAME)
```

PAGE 1-A







●4 Na+

IT 3251-23-8, Cupric nitrate 10141-05-6, Cobalt
 nitrate 10421-48-4, Ferric nitrate 13138-45-9,
 Nickel nitrate 13770-18-8, Cupric perchlorate
 34946-82-2, Cupric triflate 38465-60-0, Cupric
 tetrafluoroborate 73131-99-4
 RL: NUU (Other use, unclassified); RCT (Reactant); RACT (Reactant
 or reagent); USES (Uses)
 (materials for degrading contaminants)
RN 3251-23-8 HCAPLUS
CN Nitric acid, copper(2+) salt (8CI, 9CI) (CA INDEX NAME)

●1/2 Cu(II)

RN 10141-05-6 HCAPLUS CN Nitric acid, cobalt(2+) salt (8CI, 9CI) (CA INDEX NAME)

●1/2 Co(II)

RN 10421-48-4 HCAPLUS CN Nitric acid, iron(3+) salt (8CI, 9CI) (CA INDEX NAME)

●1/3 Fe(III)

RN 13138-45-9 HCAPLUS CN Nitric acid, nickel(2+) salt (8CI, 9CI) (CA INDEX NAME)

●1/2 Ni(II)

RN 13770-18-8 HCAPLUS CN Perchloric acid, copper(2+) salt (8CI, 9CI) (CA INDEX NAME)

●1/2 Cu(II)

RN 34946-82-2 HCAPLUS CN Methanesulfonic acid, trifluoro-, copper(2+) salt (9CI) (CA INDEX NAME)

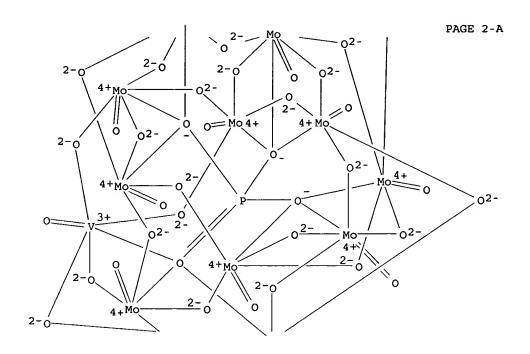
●1/2 Cu(II)

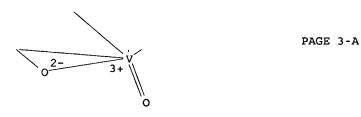
RN 38465-60-0 HCAPLUS CN Borate(1-), tetrafluoro-, copper(2+) (2:1) (9CI) (CA INDEX NAME)

●1/2 Cu(II) 2+

RN 73131-99-4 HCAPLUS
CN Vanadate(5-), (heptadeca-μ-oxodecaoxodecamolybdate)hepta-μoxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ
':κΟ':κΟ':κΟ'':κΟ'':κΟ'''
:κΟ''':κΟ''']]di-, pentasodium (9CI) (CA INDEX NAME)

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





•5 Na+

IT 59858-44-5P 134360-58-0P

RL: NUU (Other use, unclassified); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)

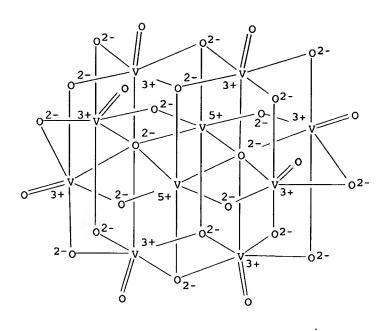
(materials for degrading contaminants)

RN 59858-44-5 HCAPLUS

CN 1-Butanaminium, N,N,N-tributyl-, tetradeca-μ-oxo-tetra-μ3oxodi-μ6-oxooctaoxodecavanadate(6-) (6:1) (9CI) (CA INDEX NAME)

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CRN 12397-12-5 CMF 028 V10 CCI CCS



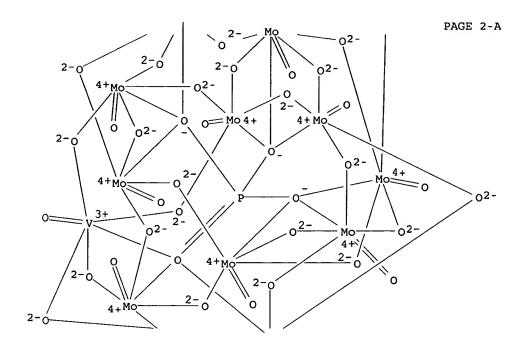
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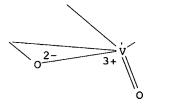
CRN 10549-76-5 CMF C16 H36 N

CRN 58071-93-5 CMF Mo10 O40 P V2 CCI CCS

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

Les Henderson





CM 2

CRN 10549-76-5 CMF C16 H36 N

IC ICM A61L

CC 4-3 (Toxicology)

ST chem warfare agent decontamination; oxometalate cationic silica chem warfare agent decontamination; copper salt oxometalate cationic silica chem warfare agent decontamination

IT Infection

(anthrax; materials for degrading contaminants)

IT Biological warfare agents
Chemical warfare agents
Decontamination

```
(materials for degrading contaminants)
ΙT
     Heteropoly acids
     RL: NUU (Other use, unclassified); USES (Uses)
        (materials for degrading contaminants)
IT
     Aldehydes, reactions
     Halogen compounds
     RL: RCT (Reactant); REM (Removal or disposal); PROC (Process);
     RACT (Reactant or reagent)
        (materials for degrading contaminants)
     Nitrates, reactions
     RL: NUU (Other use, unclassified); RCT (Reactant); RACT (Reactant
     or reagent); USES (Uses)
        (transition metal; materials for degrading contaminants
IT
     625455-59-6
                  625455-61-0 625830-47-9
                                              625830-48-0
     625830-49-1
                  625830-52-6
     RL: NUU (Other use, unclassified); RCT (Reactant); RACT (Reactant
     or reagent); USES (Uses)
        (cationic catalyst support; materials for degrading
        contaminants)
IT
     173358-70-8, Bindzil CAT
     RL: NUU (Other use, unclassified); USES (Uses)
        (materials for degrading contaminants)
     3251-23-8, Cupric nitrate 10141-05-6, Cobalt
     nitrate 10421-48-4, Ferric nitrate 12200-88-3
     13138-45-9, Nickel nitrate 13770-18-8, Cupric
     perchlorate 34946-82-2, Cupric triflate
     38465-60-0, Cupric tetrafluoroborate 73131-99-4
     625830-46-8
                  625830-51-5
     RL: NUU (Other use, unclassified); RCT (Reactant); RACT (Reactant
     or reagent); USES (Uses)
        (materials for degrading contaminants)
TT
     59858-44-5P 134360-58-0P
                               194925-14-9P
     RL: NUU (Other use, unclassified); RCT (Reactant); SPN (Synthetic
     preparation); PREP (Preparation); RACT (Reactant or reagent); USES
     (Uses)
        (materials for degrading contaminants)
IT
     1941-26-0, Tetraethylammonium nitrate
                                             1941-27-1,
     Tetrabutylammonium nitrate
     RL: NUU (Other use, unclassified); REM (Removal or disposal); PROC
     (Process); USES (Uses)
        (materials for degrading contaminants)
IT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (materials for degrading contaminants)
     50-00-0, Formaldehyde, reactions 74-93-1, Methyl mercaptan, reactions 75-07-0, Acetaldehyde, reactions 75-18-3, Dimethyl
IT
     sulfide 75-44-5, Phosgene 75-50-3, Trimethylamine, reactions
     79-09-4, Propionic acid, reactions 100-42-5, Styrene, reactions
     107-44-8, Sarin 107-92-6, n-Butyric acid, reactions 109-52-4,
     n-Valeric acid, reactions 110-01-0, Tetrahydrothiophene
     110-81-6, Diethyl disulfide 110-86-1, Pyridine, reactions
     352-93-2, Diethyl sulfide 503-74-2, Isovaleric acid
     Dimethyl disulfide 630-08-0, Carbon monoxide, reactions
     693-07-2, 2-Chloroethyl ethyl sulfide 7440-38-2D, Arsenic,
               7664-41-7, Ammonia, reactions 7704-34-9D, Sulfur,
     compds.
               7727-37-9D, Nitrogen, compds.
                                                7783-06-4, Hydrogen
     compds.
     sulfide, reactions
     RL: RCT (Reactant); REM (Removal or disposal); PROC (Process);
     RACT (Reactant or reagent)
        (materials for degrading contaminants)
L114 ANSWER 22 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
                         2003:796298 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         139:293699
                         Liquid cleaning compositions
TITLE:
```

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containing proteolytic enzyme and
```

polyoxometalate

INVENTOR(S): Adriaanse, Arend Jan; Van Dijk, Willem Robert;

Hage, Ronald

PATENT ASSIGNEE(S): Unilever Home & Personal Care, USA, Division

of Conopco, Inc., USA U.S. Pat. Appl. Publ., 19 pp. SOURCE:

CODEN: USXXCO

DOCUMENT TYPE:

Patent English

LANGUAGE: FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

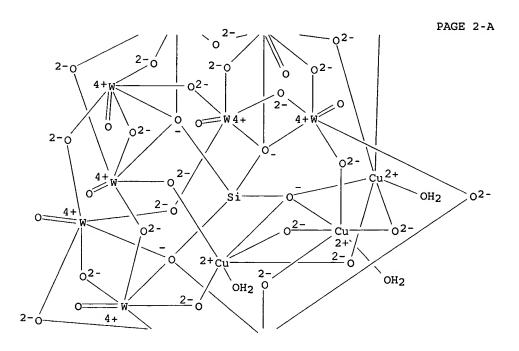
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US 2003191040	A1 2003	31009 US 2003-397413	2003
CA 2476321	AA 2003	31009 CA 2003-2476321	0326 2003
WO 2003083030	A1 2003	31009 WO 2003-EP1398	0211
			2003 0211
CH, CN, C GB, GD, G KP, KR, K MN, MW, M	O, CR, CU, CZ, E, GH, GM, HR, Z, LC, LK, LR, K, MZ, NO, NZ, K, SL, TJ, TM,	AZ, BA, BB, BG, BR, BY, BZ, DE, DK, DM, DZ, EC, EE, ES, HU, ID, IL, IN, IS, JP, KI, LS, LT, LU, LV, MA, MD, MC, OM, PH, PL, PT, RO, RU, SC, TN, TR, TT, TZ, UA, UG, UZ	S, FI, E, KG, G, MK, C, SD,
RW: GH, GM, K AZ, BY, K DE, DK, E PT, SE, S	E, LS, MW, MZ, G, KZ, MD, RU, E, ES, FI, FR,	SD, SL, SZ, TZ, UG, ZM, ZW, TJ, TM, AT, BE, BG, CH, CY, GB, GR, HU, IE, IT, LU, MC, BJ, CF, CG, CI, CM, GA, GN	Y, CZ, C, NL,
AU 2003205759		31013 AU 2003-205759	2003 0211
EP 1487954	A1 2004	EP 2003-702631	2003 0211
	E, SI, LT, LV,	FR, GB, GR, IT, LI, LU, NI FI, RO, MK, CY, AL, TR, BO	L, SE,
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		WO 2003-EP1398	W 2003 0211

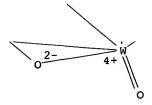
AB An aqueous liquid cleaning composition comprises a proteolytic enzyme and a primary stabilizer such as sodium tetraborate, as well as a polyoxometalate as a secondary enzyme stabilizer, such as (Me3NH) 4 (NbO2) PW11O39.

ΙT 165275-44-5 608526-64-3

RL: TEM (Technical or engineered material use); USES (Uses) (secondary enzyme stabilizer; liquid cleaning compns. containing proteolytic enzyme and polyoxometalate)



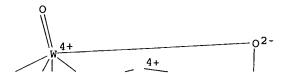


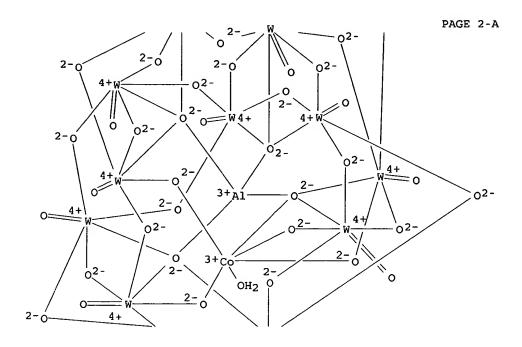


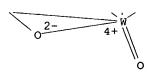
●10 K+

RN 608526-64-3 HCAPLUS CN

Tungstate(6-), aluminate(aquacobaltate)tetracosa- μ -oxotetra- μ 4-oxoundecaoxoundeca-, hexasodium (9CI) (CA INDEX NAME)





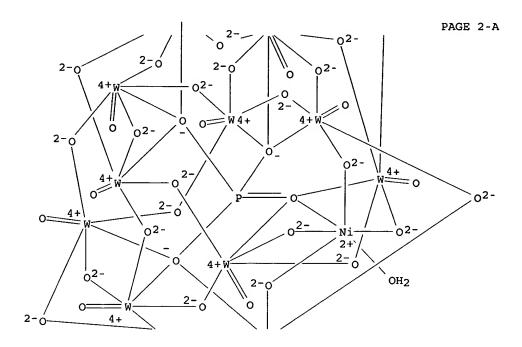


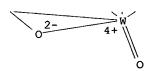
●6 Na+

IC ICM C11D003-386 INCL 510267000; 510392000; 510530000 46-5 (Surface Active Agents and Detergents) sodium tetraborate proteolytic enzyme stabilizer detergent; ST polyoxometalate enzyme stabilizer detergent IT Detergents (bleaching; liquid cleaning compns. containing proteolytic enzyme and polyoxometalate) IT Detergents (laundry, enzyme-containing; liquid cleaning compns . containing proteolytic enzyme and polyoxometalate) IT (liquid; liquid cleaning compns. containing proteolytic enzyme and polyoxometalate) IT Heteropoly acids RL: TEM (Technical or engineered material use); USES (Uses) (secondary enzyme stabilizer; liquid cleaning compns. containing proteolytic enzyme and polyoxometalate) IT 37259-58-8D, Serine protease, modified RL: TEM (Technical or engineered material use); USES (Uses) (bacterial; liquid cleaning compns. containing proteolytic enzyme and polyoxometalate) IT 1330-43-4, Sodium tetraborate 7775-19-1, Sodium metaborate

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10043-35-3, Boric acid, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (enzyme stabilizer; liquid cleaning compns.
        containing proteolytic enzyme and polyoxometalate)
IT
     9014-01-1, Subtilisin
     RL: NUU (Other use, unclassified); USES (Uses)
        (liquid cleaning compns. containing proteolytic
        enzyme and polyoxometalate)
     9001-92-7, Proteolytic enzyme
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (protease; liquid cleaning compns. containing
        proteolytic enzyme and polyoxometalate)
                               172304-20-0
IT
     110717-67-4 165275-44-5
                                            273201-44-8
     608526-64-3
                  608526-65-4
     RL: TEM (Technical or engineered material use); USES (Uses)
        (secondary enzyme stabilizer; liquid cleaning
        compns. containing proteolytic enzyme and
        polyoxometalate)
L114 ANSWER 23 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2003:574606 HCAPLUS
DOCUMENT NUMBER:
                         140:67483
TITLE:
                         Preparation, characterization, and
                         photocatalytic activity of POM-APS-silica
                         hybrid catalysts
AUTHOR (S):
                         Li, Li; Guo, Yi-xing; Yang, Yu; Zhou, Ping;
                         Jiang, Chun-jie
CORPORATE SOURCE:
                         Faculty of Chemistry and Chemical Engineering,
                         Qiqihar University, Qiqihar, 16100, Peop. Rep.
                         China
SOURCE:
                         Fenzi Kexue Xuebao (2003), 19(1), 33-39
                         CODEN: JMOSE7; ISSN: 1000-9035
PUBLISHER:
                         Dongbei Shifan Daxue Xueshu Jikanshe
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         Chinese
ΔR
     Amine-functionalized mesoporous silica materials
     impregnated with transition- metal-monosubstituted
     polyoxometalate cluster, K5[Ni(H2O)PW11O39] (PW11Ni), were
     prepared by coordination of nickel centers in the cluster with the
     amine surface groups in silica supports. XRD, UV/DRS,FT - IR, ICP
     - AES, Elemental anal. were used to characterize the structure and
     composition of the composite, and the photocatalytic
     activity of the composite was studied through
     photocatalytic degradation of dye Rhodamine B(RB).
                                                         The exptl. results
     indicated that the photocatalytic activity of the
     composite was higher than that of the direct photolysis
     and the pure PW11Ni in the homogeneous system.
     Moreover, this kind of catalyst was insol., and it could be
     reused.
TΥ
     37194-75-5
     RL: CAT (Catalyst use); USES (Uses)
        (polyoxometalate-amine-functionalized silica hybrid
        photocatalysts)
RN
     37194-75-5 HCAPLUS
CN
     Tungstate (5-), (aquanickelate) tetracosa-μ-oxoundecaoxo [μ12-
     [phosphato(3-)-\kappa0:\kappa0:\kappa0:\kappa0':\kappa0':.kap
     pa.0'κ0'':κ0'':κ0'':κ0''':κ0''':.kap
     pa.O''']]undeca-, pentapotassium (9CI) (CA INDEX NAME)
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^{*} STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





●5 K+

CC 74-1 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST polyoxometalate amine functionalized silica hybrid photocatalyst

IT Photolysis catalysts

(polyoxometalate-amine-functionalized silica hybrid photocatalysts)

IT Heteropoly acids

RL: CAT (Catalyst use); USES (Uses)

(polyoxometalate-amine-functionalized silica hybrid photocatalysts)

IT 81-88-9, Rhodamine B

RL: RCT (Reactant); RACT (Reactant or reagent) (decomposition catalyzed by polyoxometalate

-amine-functionalized silica hybrid photocatalysts)

IT 919-30-2D, silica derivs. 7631-86-9, Silica, uses 37194-75-5

RL: CAT (Catalyst use); USES (Uses)

(polyoxometalate-amine-functionalized silica hybrid photocatalysts)

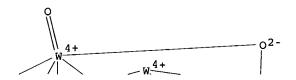
L114 ANSWER 24 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

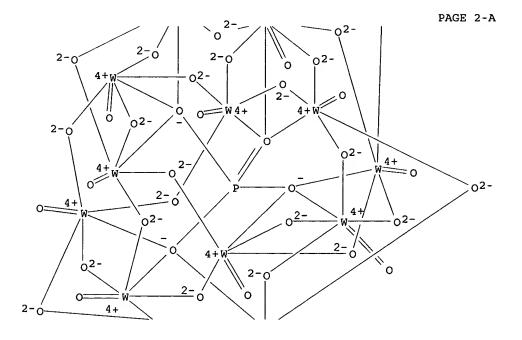
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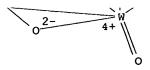
DOCUMENT NUMBER: 140:69574

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Synthesis of Heteropoly Acids and Their Salts
                         Using Mechanochemical Activation
                         Molchanov, V. V.; Maksimov, G. M.;
AUTHOR(S):
                         Maksimovskaya, R. I.; Goidin, V. V.; Buyanov,
                         R. A.
CORPORATE SOURCE:
                         Siberian Division, Boreskov Institute of
                         Catalysis, Russian Academy of Sciences,
                         Novosibirsk, 630090, Russia
SOURCE:
                         Inorganic Materials (Translation of
                         Neorganicheskie Materialy) (2003), 39(7),
                         687-693
                         CODEN: INOMAF; ISSN: 0020-1685
PUBLISHER:
                         MAIK Nauka/Interperiodica Publishing
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     A method is proposed for the synthesis of heteropoly acids from
     oxides of Mo, W, and V via mechanochem. activation. The
     fundamental principles of this approach to the synthesis of
     heteropoly acids containing different ligands and heteroatoms are
     formulated. The new V2O5 nMoO3 compds. synthesized
     in this work are highly reactive with H3PO4, which is due to the
     unsatd. coordination of the V cations and the low structural
     perfection of these compds. The application area of the proposed
     method is established. It appears to be most effective in the
     synthesis of phosphomolybdovanadic and phosphomolybdic heteropoly
     acids. In some instances, heteropoly acids can be prepared by
     solid-state reactions, which makes it possible to use
     V205 \cdot nMoO3 compds. with n \le 6 as starting reagents.
     The method has a number of important advantages: the process is
     waste-free, requires a shorter synthesis time and involves a
     smaller number of steps as compared to the existing processes,
    affords an increased yield of heteropoly acids, and involves no
     explosion- or fire-hazardous steps.
ΙT
     1343-93-7P, 12-Tungstophosphoric acid 11074-20-7P
     12026-57-2P, 12-Molybdophosphoric acid 12027-12-2P
     , 12-Molybdosilicic acid 12027-38-2P, 12-Tungstosilicic
    acid 12293-15-1P 12293-21-9P
     12293-24-2P 12398-73-1P 54327-43-4P
     92627-47-9P 92627-49-1P 92627-50-4P
     92627-51-5P 174281-83-5P 638165-41-0P
     638165-43-2P 638989-95-4P
    RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation via mechanochem. activation.)
    1343-93-7 HCAPLUS
RN
CN
    Tungstate (3-), tetracosa-\mu-oxododecaoxo [\mu12-[phosphato(3-)-
     κ0:κ0:κ0:κ0':κ0':κ0':κ0'
     ':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec
     a-, trihydrogen (9CI) (CA INDEX NAME)
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TITLE:

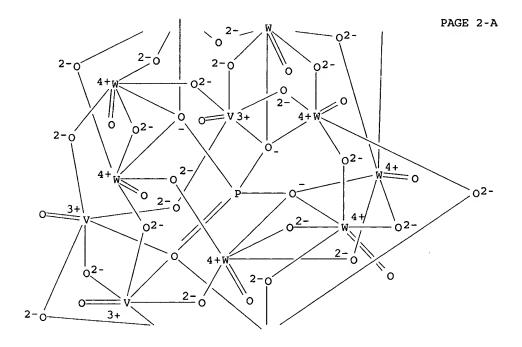






●3 H+

RN 11074-20-7 HCAPLUS
CN Vanadate(7-), (dodeca-μ-oxooctaoxooctatungstate)dodeca-μoxotetraoxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:λapp
a.O':κΟ':κΟ':κΟ'':κΟ'':κΟ
''':κΟ''':κΟ''']]tetra-, heptahydrogen (9CI) (CA
INDEX NAME)

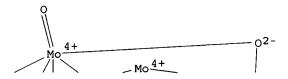


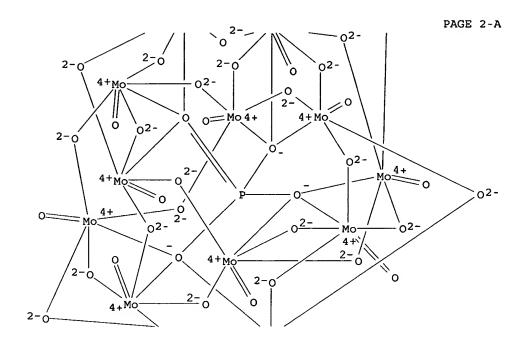
0²⁻ 3+

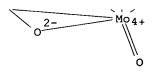
PAGE 3-A

●7 111

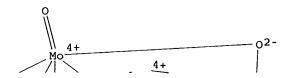
RN 12026-57-2 HCAPLUS CN Molybdate(3-), tetracosa- μ -oxododecaoxo[μ 12-[phosphato(3-)- κ 0: κ 0: κ 0: κ 0': κ 0': κ 0': κ 0'': κ 0''': κ 0'': κ 0''': κ 0'': κ 0

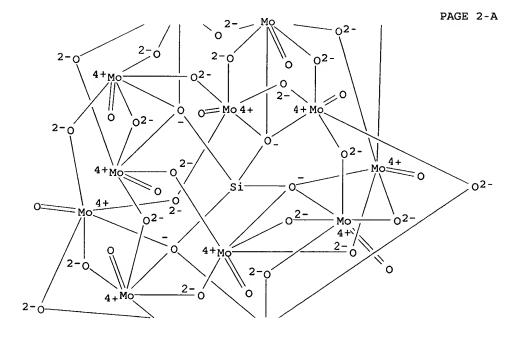


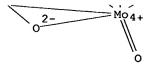




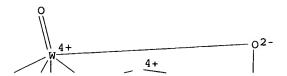
●3 H+

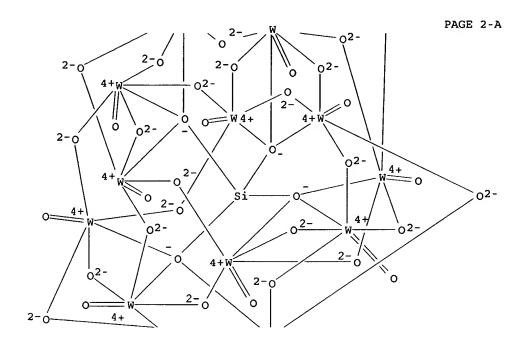






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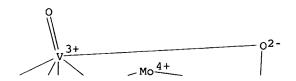


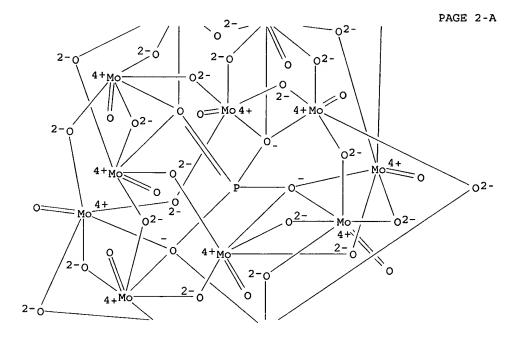


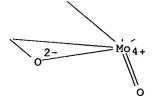


●4 H+

RN 12293-15-1 HCAPLUS
CN Vanadate(4-), (eicosa- μ -oxoundecaoxoundecamolybdate)tetra- μ -oxooxo[μ 12-[phosphato(3-)- κ 0: κ 0: κ 0: κ 0: κ 0: κ 0': κ 0': κ 0'': κ 0''

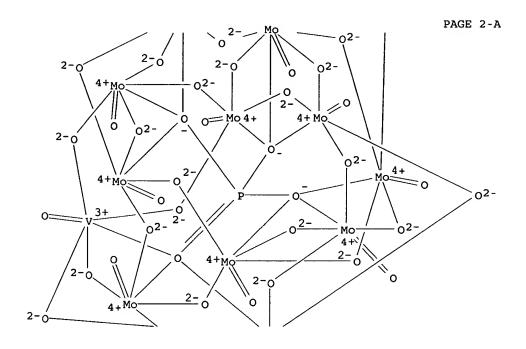


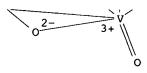




●4 H

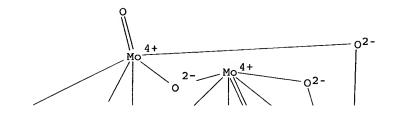


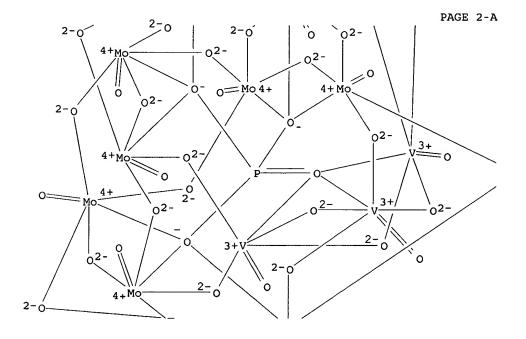




●5 H+

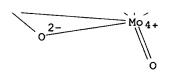
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PAGE 2-B

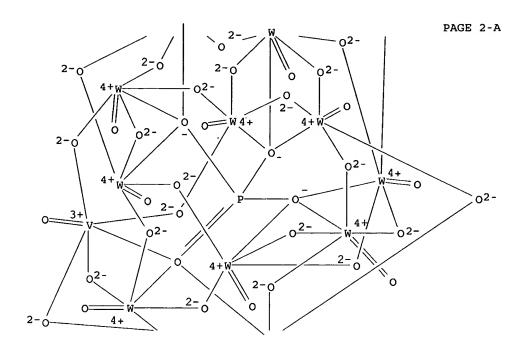
>02-

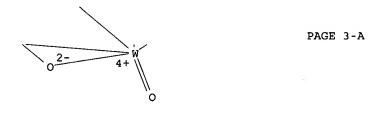


PAGE 3-A

●6 H+

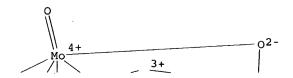
RN 12398-73-1 HCAPLUS
CN Vanadate(4-), (eicosa-μ-oxoundecaoxoundecatungstate)tetra-μoxooxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ':
κΟ':κΟ':κΟ'':κΟ'':κΟ''':
kappa.O''':κΟ''']]-, tetrahydrogen (9CI) (CA INDEX NAME)

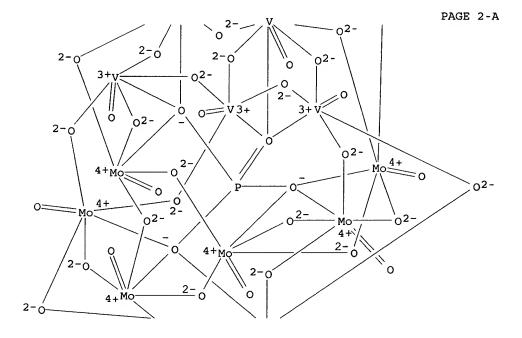


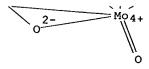


●4 H+

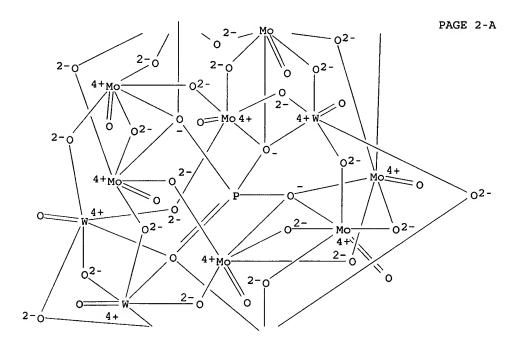
PAGE 1-A

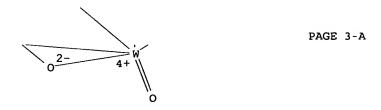


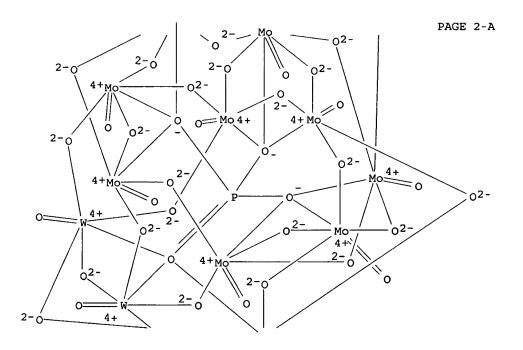


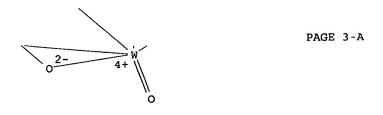


●7 H+

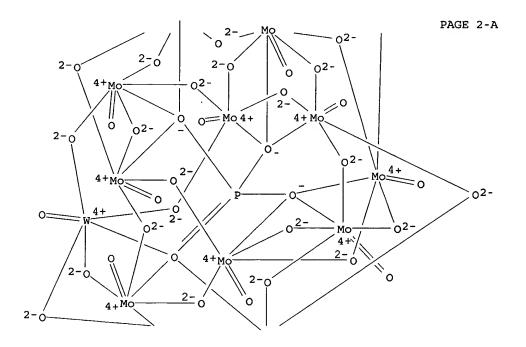






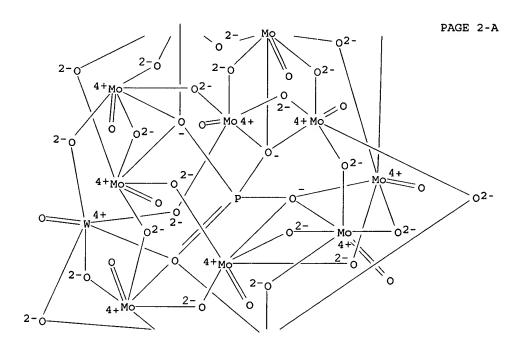


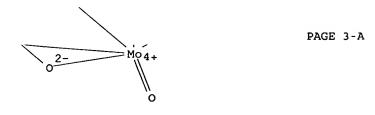
RN 92627-50-4 HCAPLUS
CN Tungstate(3-), (heptadeca-μ-oxodecaoxodecamolybdate)hepta-μoxodioxo[μ12-[phosphato(3-)-κ0:κ0:κ0
':κ0':κ0':κ0'':κ0'':κ0'':κ0'''
:κ0''':κ0''']]di-, trihydrogen (9CI) (CA INDEX NAME)



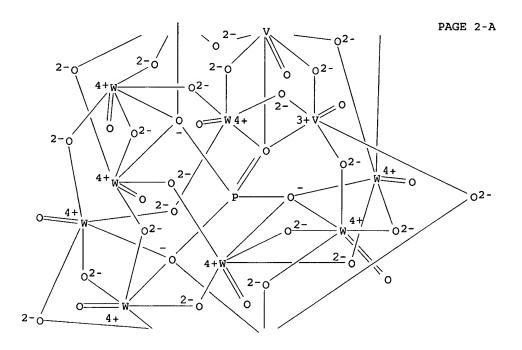


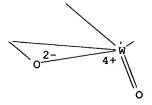
RN 92627-51-5 HCAPLUS
CN Tungstate(3-), (eicosa-μ-oxoundecaoxoundecamolybdate)tetra-μoxooxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ:
κΟ':κΟ':κΟ'':κΟ'':κΟ''':
kappa.O''':κΟ''']]-, trihydrogen (9CI) (CA INDEX NAME)



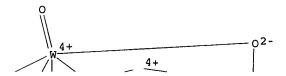


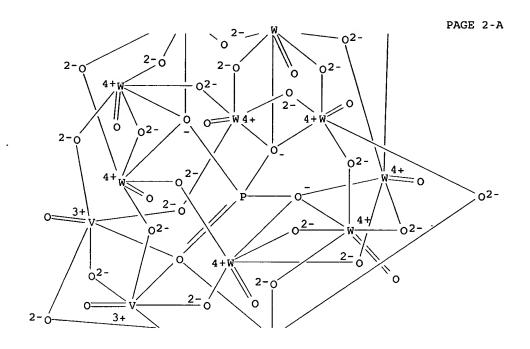
RN 174281-83-5 HCAPLUS
CN Vanadate(5-), (heptadeca-μ-oxodecaoxodecatungstate)hepta-μoxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ
':κΟ':κΟ':κΟ'':κΟ'':κΟ'''
:κΟ''':κΟ''']]di-, pentahydrogen (9CI) (CA INDEX
NAME)

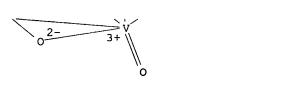




●5 H+





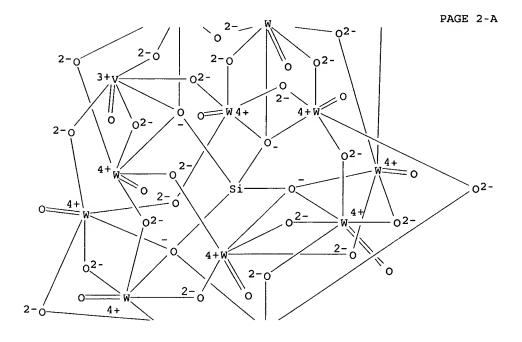


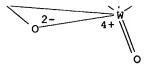
●6 H+

RN 638165-43-2 HCAPLUS
CN Vanadate(6-), (heptadeca-μ-oxodecaoxodecatungstate) [μ12[orthosilicato(4-)-κ0:κ0:κ0':κ0':
κ0':κ0'':κ0'':κ0''':κ0'''
:κ0'''] hepta-μ-oxodioxodi-, hexahydrogen (9CI) (CA
INDEX NAME)

PAGE 1-A

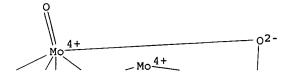


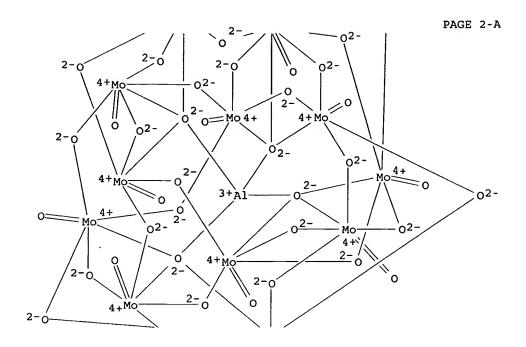


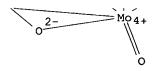


●6 H⁴

RN 638989-95-4 HCAPLUS
CN Molybdate(5-), aluminatetetracosa-µ-oxotetra-µ4oxododecaoxododeca-, pentahydrogen (9CI) (CA INDEX NAME)







DOCUMENT NUMBER:

TITLE:

AUTHOR(S):

PAGE 3-A

●5 H+

```
78-7 (Inorganic Chemicals and Reactions)
CC
     1343-93-7P, 12-Tungstophosphoric acid 11074-20-7P
TT
     12026-57-2P, 12-Molybdophosphoric acid 12027-12-2P
     , 12-Molybdosilicic acid 12027-38-2P, 12-Tungstosilicic
     acid 12207-90-8P 12293-15-1P 12293-21-9P
     12293-24-2P 12398-73-1P 54327-43-4P
     76771-55-6P 92627-47-9P 92627-49-1P
     92627-50-4P 92627-51-5P 174281-83-5P
                    638165-42-1P 638165-43-2P
     638165-41-0P
     638989-95-4P
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation via mechanochem. activation.)
REFERENCE COUNT:
                               THERE ARE 27 CITED REFERENCES AVAILABLE
                         27
                               FOR THIS RECORD. ALL CITATIONS AVAILABLE
                               IN THE RE FORMAT
L114 ANSWER 25 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2003:236094 HCAPLUS
```

ambient temperature

139:89622

Les Henderson Page 146 571-272-2538

Highly selective and efficient O2/air-based oxidation of 2-chloroethyl ethyl sulfide at

Polyoxometalates on cationic silica

Okun, Nelya M.; Anderson, Travis M.;

```
Hill, Craig L.
                         Department of Chemistry, Emory University,
CORPORATE SOURCE:
                         Atlanta, GA, 30322, USA
                         Journal of Molecular Catalysis A: Chemical
SOURCE:
                         (2003), 197(1-2), 283-290
                         CODEN: JMCCF2; ISSN: 1381-1169
                         Elsevier Science B.V.
PUBLISHER:
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     Binary cupric nitrate and triflate systems catalyze the
     homogeneous air oxidation of the mustard (HD) simulant 2-chloroethyl
     Et sulfide (CEES) to the corresponding desired sulfoxide (CEESO)
     with effectively quant. selectivity in acetonitrile under ambient
     conditions. This activity is enhanced when cationic silica
     nanoparticles coated with the anionic multi-iron
    polyoxometalates (POMs) are also present. The POM-coated
     nanoparticles are prepared by treatment of aqueous suspensions of
     Bindzil CAT cationic silica nanoparticles (from Akzo Nobel) with
    aqueous solns. of the POMs, K9[(FeIII(OH2)2)3(PW9O34)2] (K94) or Na12[(FeOH2)2Fe2(P2W15O56)2] (Na125).
CC
    60-4 (Waste Treatment and Disposal)
ST
    polyoxometalate cationic silica aerobic oxidn
     chloroethyl Et sulfide; hazardous waste polyoxometalate
     cationic silica oxidn chloroethyl Et sulfide; catalytic oxidn
     mustard simulant hazardous waste polyoxometalate
     cationic silica
    Heteropoly acids
    RL: CAT (Catalyst use); MOA (Modifier or additive use); USES
        (polyoxometalates on cationic silica for highly
        selective and efficient aerobic oxidation of 2-chloroethyl Et
        sulfide at ambient temperature)
IT
    Hazardous wastes
        (treatment; polyoxometalates on cationic silica for
        highly selective and efficient aerobic oxidation of 2-chloroethyl
        Et sulfide at ambient temperature in relation to)
IT
    173358-70-8, Bindzil CAT
     RL: CAT (Catalyst use); USES (Uses)
        (cationic silica nanoparticles; polyoxometalates on
        cationic silica for highly selective and efficient aerobic
        oxidation of 2-chloroethyl Et sulfide at ambient temperature)
                                34946-82-2, Cupric triflate
TТ
    3251-23-8, Cupric nitrate
    RL: CAT (Catalyst use); USES (Uses)
        (polyoxometalates on cationic silica for highly
        selective and efficient aerobic oxidation of 2-chloroethyl Et
        sulfide at ambient temperature)
                  554449-60-4
TΨ
    130002-64-1
    RL: CAT (Catalyst use); MOA (Modifier or additive use); USES
        (polyoxometalates on cationic silica for highly
        selective and efficient aerobic oxidation of 2-chloroethyl Et
        sulfide at ambient temperature)
TΤ
    693-07-2, 2-Chloroethyl ethyl sulfide
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); POL (Pollutant); REM (Removal or disposal); OCCU
     (Occurrence); PROC (Process)
        (polyoxometalates on cationic silica for highly
        selective and efficient aerobic oxidation of 2-chloroethyl Et
        sulfide at ambient temperature)
    27998-62-5, 2-Chloroethyl ethyl sulfoxide
    RL: FMU (Formation, unclassified); FORM (Formation,
    nonpreparative)
        (polyoxometalates on cationic silica for highly
        selective and efficient aerobic oxidation of 2-chloroethyl Et
        sulfide at ambient temperature)
REFERENCE COUNT:
                               THERE ARE 37 CITED REFERENCES AVAILABLE
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FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L114 ANSWER 26 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2003:135764 HCAPLUS DOCUMENT NUMBER: 138:261154 TITLE . [(FeIII(OH2)2)3(A- α -PW9O34)2]9- on Cationic Silica Nanoparticles, a New Type of Material and Efficient Heterogeneous Catalyst for Aerobic Oxidations Okun, Nelya M.; Anderson, Travis M.; AUTHOR (S): Hill, Craig L. CORPORATE SOURCE: Department of Chemistry, Emory University, Atlanta, GA, 30322, USA SOURCE: Journal of the American Chemical Society (2003), 125(11), 3194-3195 CODEN: JACSAT; ISSN: 0002-7863 PUBLISHER: American Chemical Society DOCUMENT TYPE: Journal LANGUAGE: English Polyoxometalates (POMs) electrostatically bind to silica nanoparticles coated with cationic aluminum oxide "(Si/AlO2)n+" to form a new type of material (the anionic POMs replace Clcounterions associated with the cationic surface sites). Association of a new .apprx.D3h POM of formula [(FeIII(OH2)2)3(A- α -PW9034)2]9- (1) with the cationic nanoparticles (to form "K81/(Si/AlO2)") was studied in detail. Elemental anal., particle sizes from both laser light scattering and TEM before and after association of 1, the size of 1 from X-ray crystallog., and other methods provide mutually consistent data that indicate about 58 K8[(FeIII(OH2)2)3(A- α -PW9O34)2]- monoanions associate with the average nanoparticle (diameter of the K81/(Si/AlO2) product = .apprx.17 nm). While heterogeneity of the cationic sites and roughness of the (Si/AlO2)n+ surfaces make the associated POMs structurally nonuniform, the equivalent of .apprx.1 monolayer of 1 is present in K81/(Si/AlO2). Remarkably, while 1, the precursor (Si/AlO2)n+, and the components of 1, each alone, are inactive as catalysts for 02/air-based oxidation of sulfides or aldehydes in solution, K81/(Si/AlO2) is an active catalyst for both reactions (facile reaction with air at low temperature). CC 67-1 (Catalysis, Reaction Kinetics, and Inorganic Reaction Mechanisms) Section cross-reference(s): 23, 27 ST polyoxometalate cationic silica nanoparticle heterogeneous catalyst aerobic oxidn; sulfide oxidn polyoxometalate cationic silica nanoparticle heterogeneous catalyst; aldehyde oxidn polyoxometalate cationic silica nanoparticle heterogeneous catalyst REFERENCE COUNT: 28 THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L114 ANSWER 27 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2002:957716 HCAPLUS DOCUMENT NUMBER: 138:229150 TITLE: Sonochemical preparation of photochromic nanocomposite thin film based on polyoxometalates well dispersed in polyacrylamide AUTHOR (S): Feng, Wei; Zhang, Tie Rui; Liu, Yan; Lu, Ran; Zhao, Ying Ying; Li, Tie Jin; Yao, Jian Nian

CORPORATE SOURCE:

SOURCE:

Les Henderson Page 148 571-272-2538

Department of Chemistry, Jilin University, Chang Chun, 130023, Peop. Rep. China Journal of Solid State Chemistry (2002),

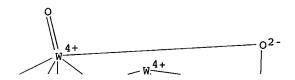
CODEN: JSSCBI; ISSN: 0022-4596

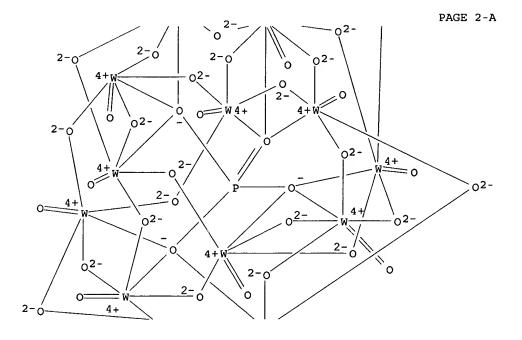
169(1), 1-5

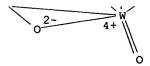
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PUBLISHER:
                        Elsevier Science
DOCUMENT TYPE:
                        Journal
LANGUAGE:
                        English
     Novel photochromic nanocomposite thin film containing phosphotungstic
     acid entrapped in polyacrylamide was prepared using ultrasound
     technique. TEM image revealed that the polyoxometalates
     nanoparticles with narrow size distribution were finely dispersed
     in polymer matrix. IR spectra showed that the Keggin geometry of
     polyoxometalates was still preserved inside the
     composite film and strong Coulombic interaction was built
     between heteropolyoxometalates and polyacrylamide via hydrogen
     bond. It was found that the thermal stability of the hybrid film
     was lower than that of pure polymer but the film was
     stable enough for photochromic application from the TG-DTA curves.
     The transparent film changed from colorless to blue under UV
     irradiation and showed reversible photochromism. The bleaching
     process occurred when the film was in contact with air or O2 in
     the dark.
               The photoreduced process was in accordance with radical
    mechanism.
IT
    12534-78-0
     RL: CPS (Chemical process); FMU (Formation, unclassified); PEP
     (Physical, engineering or chemical process); PRP (Properties);
     FORM (Formation, nonpreparative); PROC (Process)
        (sonochem. preparation of photochromic nanocomposite thin film based
        on polyoxometalates dispersed in polyacrylamide)
RN
     12534-78-0 HCAPLUS
CN
     Tungstate (4-), tetracosa-\mu-oxododecaoxo [\mu12-[phosphato(3-)-
     INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
    1343-93-7, Phosphotungstic acid
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PRP (Properties); RCT (Reactant); PROC (Process); RACT
     (Reactant or reagent)
        (sonochem. preparation of photochromic nanocomposite thin film based
       on polyoxometalates dispersed in polyacrylamide)
    1343-93-7 HCAPLUS
RN
CN
    Tungstate (3-), tetracosa-\mu-oxododecaoxo [\mu12-[phosphato(3-)-
    κ0:κ0:κ0:κ0':κ0':κ0':κ0'
```

':κΟ'':κΟ'':κΟ''':κΟ''':κΟ''']]dodec a-, trihydrogen (9CI) (CA INDEX NAME)

PAGE 1-A







●3 H+

CC 74-9 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

IT Sound and Ultrasound

(preparation of photochromic nanocomposite thin film based on polyoxometalate and polyacrylamide using)

IT Hydrogen bond

Nanocomposites

Photochromic materials

(sonochem. preparation of photochromic nanocomposite thin film based on polyoxometalates dispersed in polyacrylamide)

IT 12534-78-0

RL: CPS (Chemical process); FMU (Formation, unclassified); PEP (Physical, engineering or chemical process); PRP (Properties); FORM (Formation, nonpreparative); PROC (Process)

(sonochem. preparation of photochromic nanocomposite thin film based on polyoxometalates dispersed in polyacrylamide)

IT 1343-93-7, Phosphotungstic acid 9003-05-8,

Polyacrylamide

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PRP (Properties); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)

(sonochem. preparation of photochromic nanocomposite thin film based on polyoxometalates dispersed in polyacrylamide)

REFERENCE COUNT:

THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L114 ANSWER 28 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

21

ACCESSION NUMBER: 2002:916776 HCAPLUS

DOCUMENT NUMBER: 138:323871

TITLE: A novel application of mixed-valence

Keggin-type polyoxometalates as non-aqueous electrolytes in polyacenic semiconductor secondary batteries

AUTHOR(S): Wang, Xiuli; Wang, Enbo; Xie, Demin; Zhang,

Xiyan; Hu, Changwen; Xu, Lin

CORPORATE SOURCE: Institute of Polyoxometalate Chemistry,

Department of Chemistry, Northeast Normal University, Changchun, 130024, Peop. Rep.

China

SOURCE: Solid State Ionics (2003), 156(1,2), 71-78

CODEN: SSIOD3; ISSN: 0167-2738

PUBLISHER: Elsevier Science B.V.

DOCUMENT TYPE: Journal LANGUAGE: English

AB Mixed-valence Keggin-type lithium

polyoxometalates (POMs) were used as the electrolytes of polyacenic semiconductor (PAS) secondary batteries substituting for LiClO4 for the first time. The discharging, cycle and self-discharging properties of these PAS/Li secondary batteries and the effect of c.d. and temperature on the properties of the batteries have been investigated. The results indicate not only that the lithium POMs can overcome the disadvantages of LiClO4,

which is apt to explode when heated or rammed, but also that some of the PAS/Li secondary batteries assembled with the novel electrolytes have larger capacity and smaller self-discharging than that assembled with LiClO4. Therefore, it is believed that Keggin-type mixed-valence lithium POMs are novel and better electrolytes of PAS secondary batteries and exhibit promising practical application.

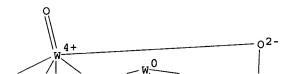
IT 514202-37-0 514202-38-1 514202-49-4

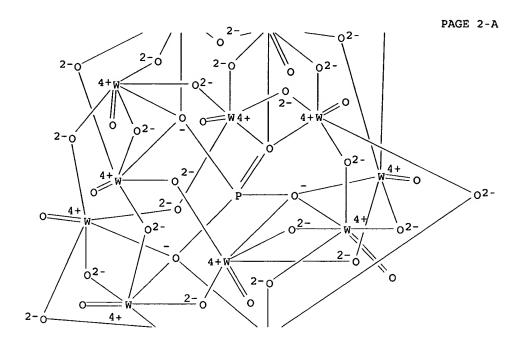
RL: DEV (Device component use); PRP (Properties); USES (Uses) (electrolytes; novel application of mixed-valence Keggin-type polyoxometalates as non-aqueous electrolytes in polyacenic semiconductor secondary batteries)

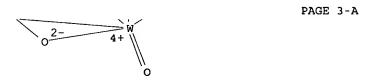
RN 514202-37-0 HCAPLUS

CN Tungstate(7-), tetracosa- μ -oxododecaoxo[μ 12-[phosphato(3-)- κ 0: κ 0: κ 0: κ 0': κ 0': κ 0'

':κ0'':κ0'':κ0''':κ0''']]dodec a-, heptalithium (9CI) (CA INDEX NAME)



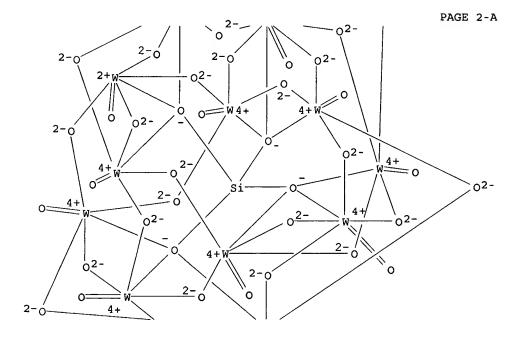


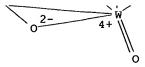


●7 Li+

PAGE 1-A

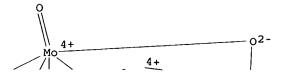






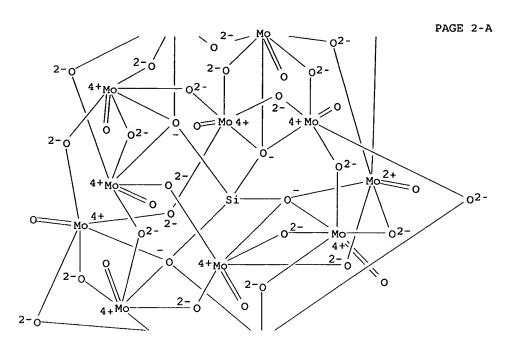
●6 Li+

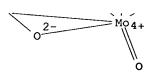
RN 514202-49-4 HCAPLUS
CN Molybdate(6-), [μ12-[orthosilicato(4-)κ0:κ0:κ0:κ0':κ0':κ0'
':κ0'':κ0'':κ0''':κ0''']]tetra
cosa-μ-oxododecaoxododeca-, hexalithium (9CI) (CA INDEX NAME)





IT





PAGE 3-A

●6 Li+

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

Secondary batteries
 (lithium; novel application of mixed-valence
 Keggin-type polyoxometalates as non-aqueous electrolytes
 in polyacenic semiconductor secondary batteries)

IT Battery electrolytes
 (novel application of mixed-valence Keggin-type
 polyoxometalates as non-aqueous electrolytes in polyacenic
 semiconductor secondary batteries)

IT Heteropoly acids
 RL: DEV (Device component use); PRP (Properties); USES (Uses)

RL: DEV (Device component use); PRP (Properties); USES (Uses) (novel application of mixed-valence Keggin-type polyoxometalates as non-aqueous electrolytes in polyacenic semiconductor secondary batteries)

514202-37-0 514202-38-1 514202-49-4
RL: DEV (Device component use); PRP (Properties); USES (Uses) (electrolytes; novel application of mixed-valence Keggin-type polyoxometalates as non-aqueous electrolytes in polyacenic semiconductor secondary batteries)



REFERENCE COUNT:

THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L114 ANSWER 29 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

30

ACCESSION NUMBER: 2002:168805 HCAPLUS

DOCUMENT NUMBER: 137:95992

TITLE: Isomerization of n-Hexane over

Silica-Supported Heteropoly Acids Promoted by

the Reduced Ce-Ni Oxides

AUTHOR(S): Kuang, Wenxing; Rives, Alain; Ben Tayeb,

Bouchta Ouled; Fournier, Michel; Hubaut,

Robert

CORPORATE SOURCE: Laboratoire de Catalyse Heterogene et

Homogene, UPRESA 8010, Universite des Sciences et Technologies de Lille, Villeneuve d'Ascq, 59655, Fr.

59655, FI

SOURCE: Journal of Colloid and Interface Science

(2002), 248(1), 123-129

CODEN: JCISA5; ISSN: 0021-9797

PUBLISHER: Academic Press

DOCUMENT TYPE: Journal LANGUAGE: English

AB The structure and catalytic properties of silica-supported heteropoly acids promoted by the reduced Ce-Ni oxides are 1st studied by using chemical anal. XRD, FT-Raman, XPS, EPR, TG, surface area measurements, and microreactor test. Silica-supported heteropoly acids have isomerization activity, but are very easy to deactivate by coke deposition. With the promotion of the reduced Ce-Ni oxides, however, the better activity and the higher selectivity to isomers could be obtained, suggesting that the reduced Ce-Ni oxides really have hydrogenating/dehydrogenating functions. Also, the presence of the reduced Ce-Ni oxides is not only beneficial for eliminating the coke deposition, but also effective for maintaining the structure of silica-supported heteropoly acids during reaction. The effect of the compn. of the mech. mixts. of silica-supported heteropoly acids and the Ce-Ni oxides on the catalytic properties was explored. (c) 2002 Academic Press.

explored. (c) 2002 Academic Press.

IT 1343-93-7, 12-Phosphotungstic acid 12027-38-2,
12-Tungstosilicic acid 12297-12-0, 12-Tungstoboric acid
RL: CAT (Catalyst use); PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

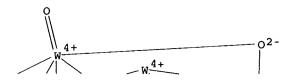
(isomerization of n-hexane over silica-supported heteropoly

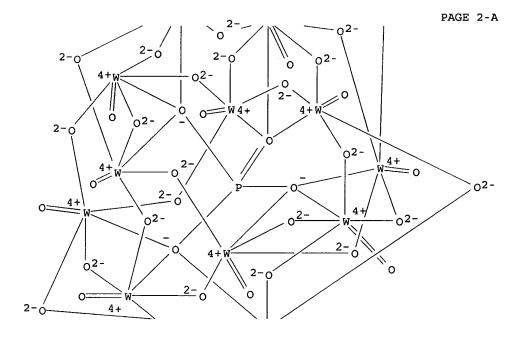
acids promoted by reduced Ce-Ni oxides)

RN 1343-93-7 HCAPLUS

CN Tungstate(3-), tetracosa-μ-oxododecaoxo[μ12-[phosphato(3-)κΟ:κΟ:κΟ:κΟ':κΟ':κΟ'

':κ0'':κ0''':κ0''':κ0''']]dodec a-, trihydrogen (9CI) (CA INDEX NAME)

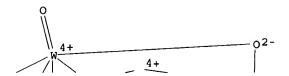


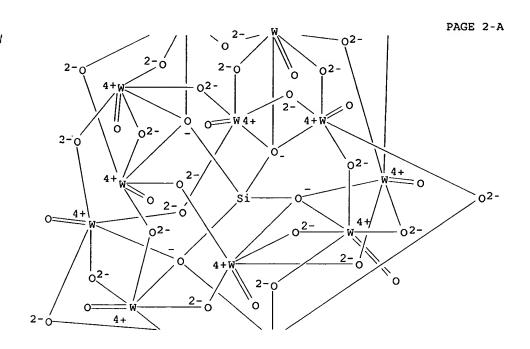


0²⁻ 4+ W

PAGE 3-A

●3 H⁴



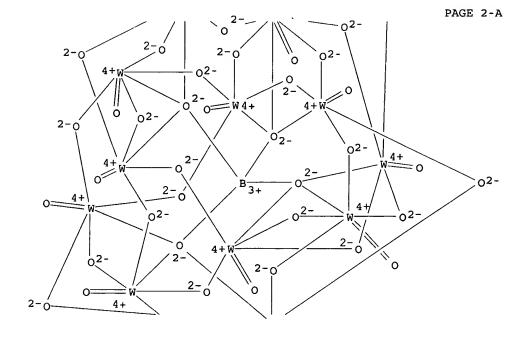


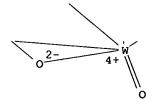


●4 H+

RN 12297-12-0 HCAPLUS CN Tungstate(5-), tetracosa- μ -oxododecaoxo[μ 12-[tetrahydroxyborato(5-)- κ 0: κ 0: κ 0: κ 0:.kapp a.0': κ 0'': κ 0''': κ 0'': κ 0''': κ







●5 H+

51-6 (Fossil Fuels, Derivatives, and Related Products) Section cross-reference(s): 45, 66 IT 1343-93-7, 12-Phosphotungstic acid 12027-38-2, 12-Tungstosilicic acid 12297-12-0, 12-Tungstoboric acid 134883-91-3, Cerium nickel oxide RL: CAT (Catalyst use); PRP (Properties); TEM (Technical or engineered material use); USES (Uses) (isomerization of n-hexane over silica-supported heteropoly acids promoted by reduced Ce-Ni oxides) REFERENCE COUNT: 53 THERE ARE 53 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L114 ANSWER 30 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

2001:795455 HCAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 136:95024

TITLE . Asymmetric Sandwich-Type Polyoxoanions.

Synthesis, Characterization, and X-ray Crystal

Structures of Diferric Complexes

[TMIIFeIII2 (P2W15056) (P2TMII2W13052)]16-, TM =

Cu or Co

AUTHOR (S): Anderson, Travis M.; Hardcastle, Kenneth I.;

Okun, Nelya; Hill, Craig L.

CORPORATE SOURCE: Department of Chemistry, Emory University,

Atlanta, GA, 30322, USA

Inorganic Chemistry (2001), 40(25), 6418-6425
CODEN: INOCAJ; ISSN: 0020-1669 SOURCE:

PUBLISHER: American Chemical Society

DOCUMENT TYPE:

Journal English

LANGUAGE:

CASREACT 136:95024

OTHER SOURCE(S): Reaction of the diferric sandwich-type polyoxometalate (NaOH2)2FeIII2(P2W15O56)216- (1) with excess aqueous Cu(II) or Co(II) yields a new type of d-electron-metal substituted polyoxometalate, [TMIIFeIII2 (P2W15O56) (P2TMII2W13052)]16-, TM = Cu (2), Co (3), resp. The structure of the Na salt of 2 (Na2), determined by single-crystal x-ray diffraction anal. (a 13.4413(9), b 21.2590(15), c 25.5207(18) Å, α 80.475(2), β 85.555(2), γ 89.563(2)°, triclinic, P.hivin.1, R1 = 5.42%, based on 43097 independent reflections), consists of a defect Fe2Cu central unit sandwiched between two different trivacant Wells-Dawson-type units, P2W15 and P2Cu2W13, where the latter unit has two octahedral Cu(II) ions substituted for two adjacent belt W(VI) atoms. The CuO5OH2 octahedron in the central unit shows pronounced Jahn-Teller distortion. A low-resolution x-ray structure of Na3 is included in the Supporting Information. UV-visible, IR, 31P NMR, cyclic voltammetric, and elemental anal. data are all consistent with the structure determined from the x-ray anal. Cyclic voltammograms of 2 and 3 exhibit multiple electron-transfer processes under ambient conditions, and

Cu or Co incorporation into the framework of 1 results in a

substantial perturbation of the electrochem. properties of the polyoxotung state framework. The Bu4N+ salts of 2 and 3 (readily prepared by metathesis) are stable and effective catalysts for the oxidation of some alkenes with high yields based on H2O2. 78-7 (Inorganic Chemicals and Reactions) Section cross-reference(s): 67, 72, 75
ENCE COUNT: 116 THERE ARE 116 CITED REFERENCES AVAILABLE REFERENCE COUNT: FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L114 ANSWER 31 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2001:721198 HCAPLUS DOCUMENT NUMBER: 136:112197 TITLE: Insulin mimetic effect of a tungstate cluster. Effect of oral administration of homo-polyoxotungstates and vanadium-substituted polyoxotungstates on blood glucose level of STZ mice AUTHOR (S): Nomiya, K.; Torii, H.; Hasegawa, T.; Nemoto, Y.; Nomura, K.; Hashino, K.; Uchida, M.; Kato, Y.; Shimizu, K.; Oda, M. CORPORATE SOURCE: Faculty of Science, Department of Materials Science, Kanagawa University, Hiratsuka, Kanagawa, 259-1293, Japan Journal of Inorganic Biochemistry (2001), SOURCE: 86(4), 657-667 CODEN: JIBIDJ; ISSN: 0162-0134 PUBLISHER: Elsevier Science Inc. DOCUMENT TYPE: Journal LANGUAGE: English Aqueous vanadate and aqueous tungstate have been known to mimic all or most of the actions of insulin in intact cell systems with respect to normalization of the blood glucose level. By carrying out oral administration in vivo expts. on the blood glucose level of streptozotocin (STZ)-induced diabetes (STZ mice), the insulin-mimetic (IM) effects of metal-oxide clusters of all-inorg. composition were examined using many types of polyoxometalates (POM) with and without vanadium substitution. Several homo-POM and vanadium-substituted POM showed hypoglycemic effects. The observed hypoglycemic effects indicated that POM with the Dawson structure { $[\alpha-P2W18062]6$ -(W-2), $[\alpha-P2W17VV062]$ 7- (V-19) and $[\alpha-1,2,3-1]$ P2W15VV3O62]9- (V-04)} are more effective than those with the Keggin structure $\{ [\alpha-PW12O40]3-(W-1), [\alpha-PW11VVO40]4-$ vanadate cluster [V10028]6- (V-15) also showed a hypoglycemic effect. 31P and 51V NMR measurements showed that the Dawson POM (W-2, V-04 and V-19) are stable in aqueous solution under the conditions used. The effect of all POM on the body weight of STZ mice was also examined The decrease in body weight after administration of W-2 was much less than for V-19, V-04 and V-15. This suggests that not only monomeric tungstate and vanadate, but also the structure factors of tungstate and vanadate clusters, can play a significant role in their biol. action. 12026-98-1 12773-19-2 37234-37-0 59519-71-0 59519-72-1 85585-36-0 93222-17-4 101144-77-8 119390-04-4 133348-27-3 133348-30-8 RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (insulin mimetic effect of a tungstate cluster. effect of oral

ΙT

RN

of STZ mice)

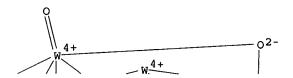
12026-98-1 HCAPLUS

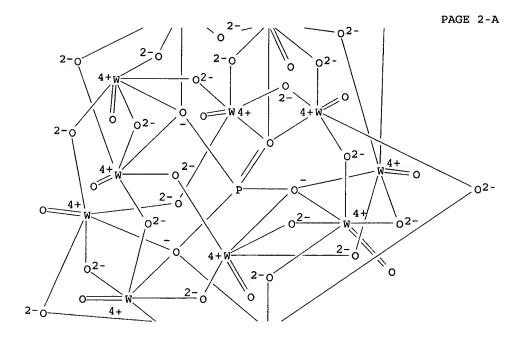
Les Henderson Page 163 571-272-2538

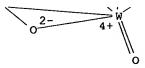
administration of homo-polyoxotungstates and

vanadium-substituted polyoxotungstates on blood glucose level

CN Tungstate(3-), tetracosa- μ -oxododecaoxo[μ 12-[phosphato(3-)- κ 0: κ 0: κ 0: κ 0': κ 0': κ 0': κ 0': κ 0'': κ 0''': κ 0'': κ 0''': κ 0'': κ 0''': κ



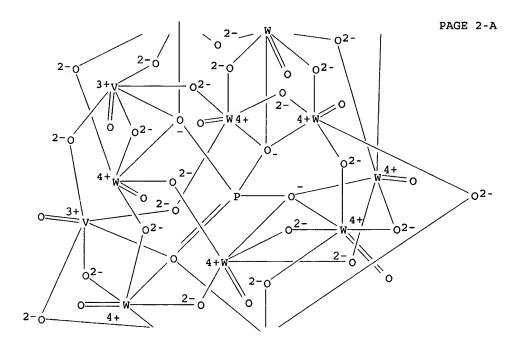




●3 Na+

RN 12773-19-2 HCAPLUS
CN Vanadate(5-), (heptadeca-μ-oxodecaoxodecatungstate)hepta-μoxodioxo[μ12-{phosphato(3-)-O:0:0':0':0':0'':0'':0'':0''
':0''']]di-, pentapotassium (9CI) (CA INDEX NAME)

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

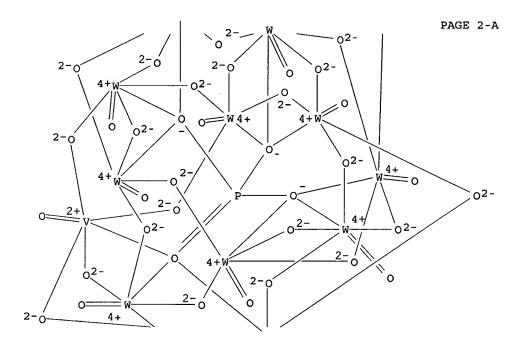


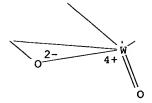


●5 K1

RN 37234-37-0 HCAPLUS
CN Vanadate(5-), (eicosa-μ-oxoundecaoxoundecatungstate)tetra-μoxooxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ:
κΟ':κΟ':κΟ'':κΟ'':κΟ''::
kappa.O''':κΟ''']]-, pentapotassium (9CI) (CA INDEX NAME)

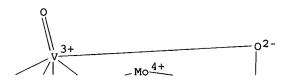
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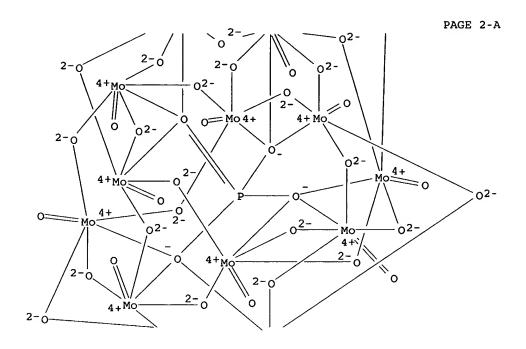


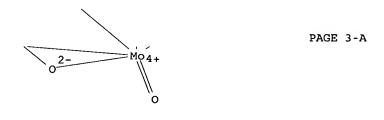


RN59519-71-0 HCAPLUS CN

Vanadate(4-), (eicosa- μ -oxoundecaoxoundecamolybdate)tetra- μ oxooxo[μ12-[phosphato(3-)-κ0:κ0:κ0:κ0':
κ0':κ0'':κ0'':κ0'':κ0''':
kappa.0''':κ0''']]-, tetrapotassium (9CI) (CA INDEX NAME)





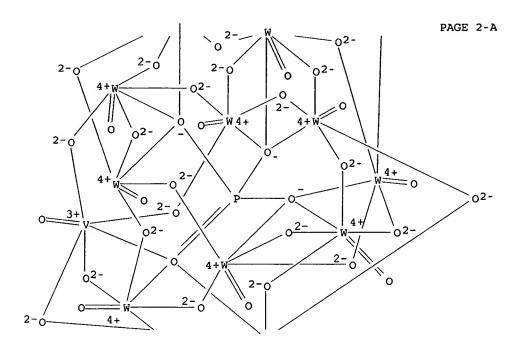


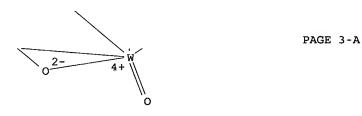
●4 K+

RN 59519-72-1 HCAPLUS
CN Vanadate(4-), (eicosa-μ-oxoundecaoxoundecatungstate)tetra-μoxooxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ':
κΟ':κΟ'':κΟ'':κΟ'':κΟ''':
kappa.O''':κΟ''']]-, tetrapotassium (9CI) (CA INDEX NAME)

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

Les Henderson Page 168 571-272-2538

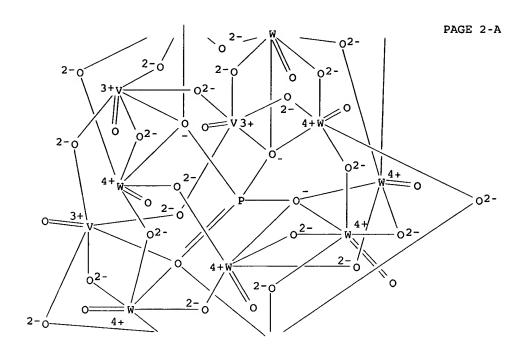


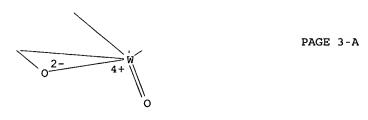


●4 K+

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85585-36-0 HCAPLUS
RN
     Vanadate(8-), [dotriaconta-\mu-oxoheptadecaoxo[\mu9-[phosphato(3-
CN
     )-κ0:κ0:κ0:κ0':κ0':κ0'':.kappa
     .O'':κO''':κO''']]heptadecatungstate]tetra-μ-
     οχοοχο [μ9-[phosphato(3-)-κ0:κ0:κ0:κ0':.
     kappa.0':κ0'':κ0'':κ0''':κ0''']]-,
     octapotassium (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN
     93222-17-4 HCAPLUS
     Vanadate(6-), nona-\mu-oxotrioxo(pentadeca-\mu-
CN
     oxononaoxononatungstate) [µ12-[phosphato(3-)-
     κ0:κ0:κ0:κ0':κ0':κ0':κ0'
     ':κ0'':κ0'':κ0''':κ0''':κ0''']]tri-,
     hexapotassium (9CI) (CA INDEX NAME)
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* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

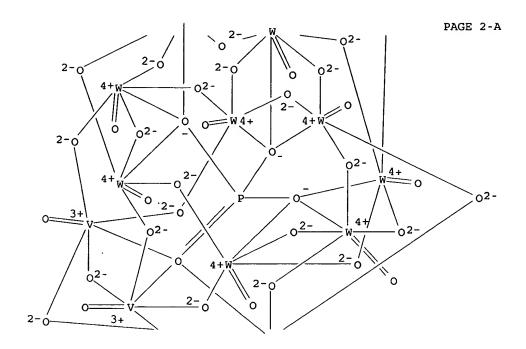


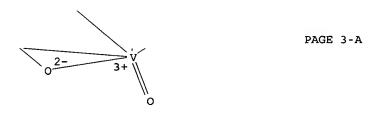


●6 K+

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RN
     101144-77-8 HCAPLUS
CN
     Vanadate(9-), [heptacosa-\mu-oxopentadecaoxo[\mu9-[phosphato(3-)-
     κ0:κ0:κ0:κ0':κ0':κ0'',κ0
     '':κΟ''':κΟ''']]pentadecatungstate]nona-μ-
     oxotrioxo[μ9-[phosphato(3-)-κ0:κ0:κ0:κ0
     ':κ0':κ0'':κ0'':κ0''':κ0''']]tri-,
     octapotassium hydrogen (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN
     119390-04-4 HCAPLUS
CN
     Vanadate(6-), nona-\mu-oxotrioxo(pentadeca-\mu-
     oxononaoxononatungstate) [µ12-[phosphato(3-)-
     κ0:κ0:κ0:κ0':κ0':κ0':κ0'
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     hexapotassium (9CI) (CA INDEX NAME)
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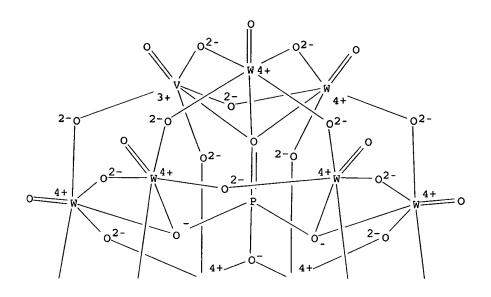
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

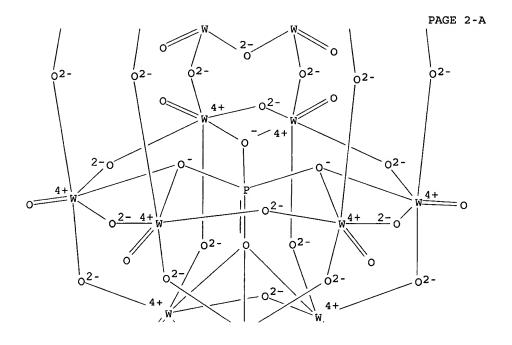




●6 K+

PAGE 1-A





RN 133348-30-8 HCAPLUS CN Vanadate(8-), $[nonacosa-\mu-oxohexadecaoxo[\mu9-[phosphato(3-)$ κ0:κ0:κ0:κ0':κ0':κ0'':κ0 $'':\kappa0''':\kappa0'''$] hexadecatungstate] hepta- μ oxodioxo $[\mu 9 - [phosphato (3-) - \kappa 0: \kappa 0: \kappa 0: \kappa 0]]$:κ0':κ0'':κ0'':κ0''':κ0''']]di-, octapotassium (9CI) (CA INDEX NAME) *** STRUCTURE DIAGRAM IS NOT AVAILABLE *** 1-3 (Pharmacology) CC IT 12026-98-1 12200-88-3 12773-19-2 13472-45-2

37234-37-0 59519-71-0 59519-72-1 85585-36-0 93222-17-4 93240-37-0 101144-77-8 119390-04-4 133348-27-3 133348-30-8

RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (insulin mimetic effect of a tungstate cluster. effect of oral administration of homo-polyoxotungstates and vanadium-substituted polyoxotungstates on blood glucose level of STZ mice)

REFERENCE COUNT:

THERE ARE 54 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L114 ANSWER 32 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2001:639652 HCAPLUS

TITLE:

AUTHOR (S):

Modifications of catalytic properties in

multi-iron sandwich-type

polyoxometalates based on iron

populations in the central tetrameric unit

Anderson, Travis M.; Zhang, Xuan; Okun,

Nelya; Chen, Qin; Hill, Craig L.

CORPORATE SOURCE:

Department of Chemistry, Emory University,

Atlanta, GA, 30322, USA

SOURCE:

Abstracts of Papers, 222nd ACS National Meeting, Chicago, IL, United States, August 26-30, 2001 (2001), INOR-465. American Chemical Society: Washington, D. C.

CODEN: 69BUZP

LANGUAGE:

DOCUMENT TYPE: Conference; Meeting Abstract English

A new diferric sandwich compound with the formula, FeIII2(NaOH2)2 (P2W15O56)2, has been prepared by the reaction of trivacant α -Na12P2W15O56 with FeCl2ullet4H2O followed by O2 oxidation in aqueous solution The complex has been fully characterized by single crystal X-ray diffraction anal. and is consistent with all IR, 31P NMR, and elemental anal. data. Unlike the tetraferric Dawson sandwich compound, the tetra-n-butylammonium salt of this complex is a highly effective catalyst for H2O2-based epoxidn. of alkenes. The substituent effects of the FeIII centers on the electrochem. and catalytic properties of the two complexes are compared.

L114 ANSWER 33 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2001:359866 HCAPLUS

DOCUMENT NUMBER:

134:362656

TITLE:

Polyoxometalate materials,

metal-containing materials, and methods of use

APPLICATION NO.

DATE

thereof

INVENTOR(S):

Hill, Craig; Xu, Ling; Rhule, Jeffrey T.;

PATENT ASSIGNEE(S):

Boring, Eric A. Emory University, USA PCT Int. Appl., 68 pp.

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CODEN: PIXXD2

KIND DATE

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Patent

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English

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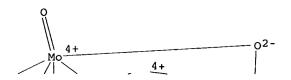
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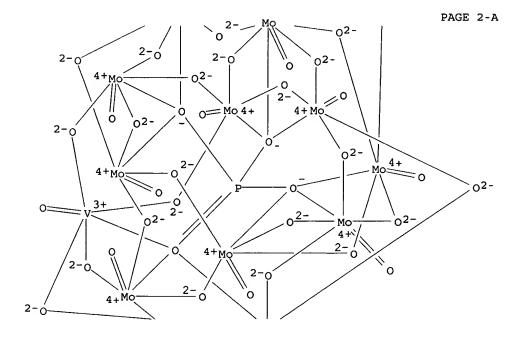
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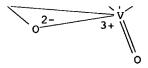
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AB The invention relates to a polyoxometalate topical

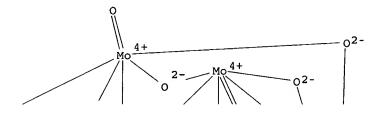
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composition for removing a contaminant from an
     environment, comprising a topical carrier and at least one
     polyoxometalate, with the proviso that the
     polyoxometalate is not H5PV2Mo10040; K5Si(H2O)MnIIIW11039;
     K4Si(H2O)MnIVW11039; or K5CoIIIW12040. The invention further
     relates to a method for removing a contaminant from an
     environment by the composition and contacting a
     polyoxometalate powder or a polyoxometalate
     coating with the environment. It further relates to a modified
     polyoxometalate, comprising the admixt. of (1) a
     polyoxometalate and (2) a cerium, a silver, a gold, a
     platinum compound, or a combination thereof. The
     invention further relates to a method for removing a
     contaminant from an environment by contacting a modified
     material comprising (1) a material and (2) a metal compound
     comprising a transition metal compound, an actinide compound, a
     lanthanide compound, or a combination thereof, wherein the
     metal compound is not a polyoxometalate. The modified
     material comprises (1) a material comprising a topical carrier, a
    powder, a coating, or a fabric, and (2) a metal compound comprising
     a transition metal compound, an actinide compound, a lanthanide
     compound, or a combination thereof, wherein the metal
     compound is not a polyoxometalate. The invention further
     relates to an article comprising the modified material.
    12293-21-9 12293-24-2 67724-86-1
    73131-99-4 73132-07-7 122795-31-7
     170663-07-7 187289-60-7 273201-47-1
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RN
    12293-21-9 HCAPLUS
    Vanadate \, (5\text{--}) \, , \quad (heptadeca-\mu-oxodecaoxodecamolybdate) \, hepta-\mu-
CN
    oxodioxo [μ12-[phosphato (3-)-κ0:κ0:κ0:κ0
     ':κ0':κ0':κ0'':κ0'':κ0'':κ0''
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    NAME)
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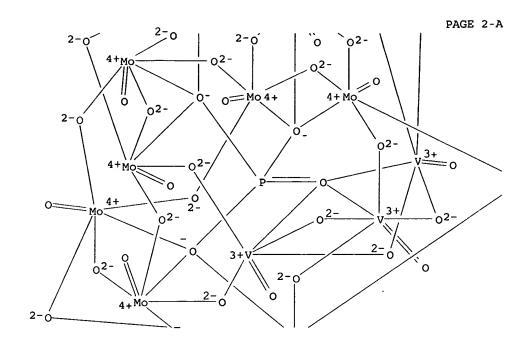




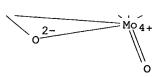


●5 114





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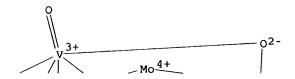


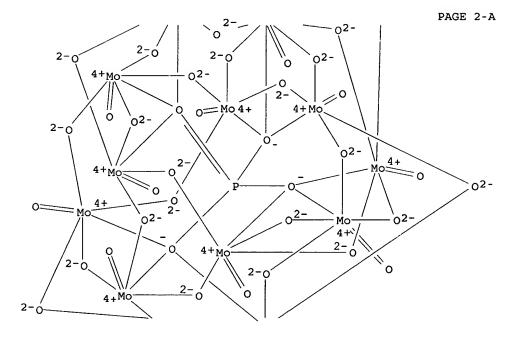
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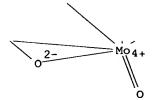
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PAGE 1-A

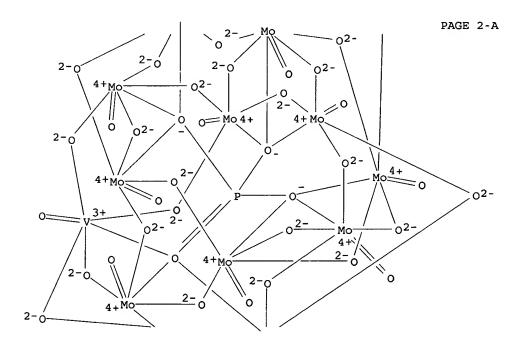


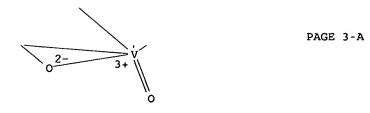




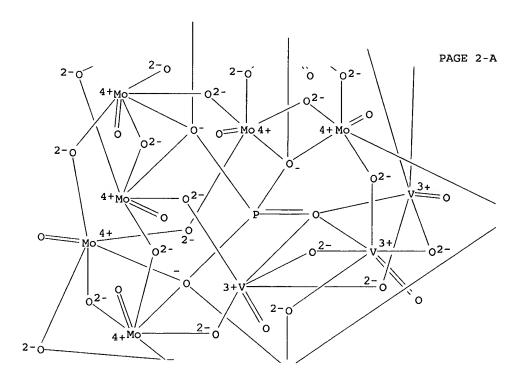
●4 Na+

RN 73131-99-4 HCAPLUS
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':κΟ':κΟ':κΟ'':κΟ'':κΟ'''
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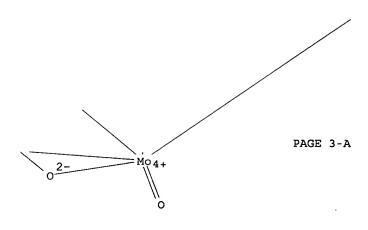


●5 Na+



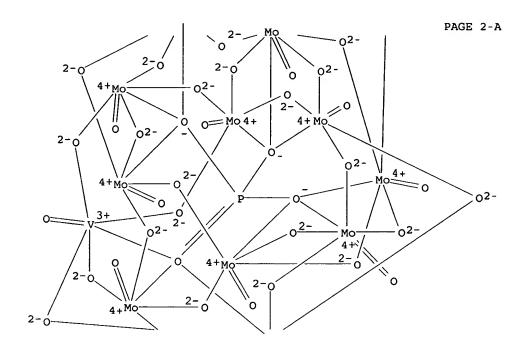
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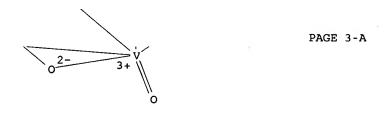
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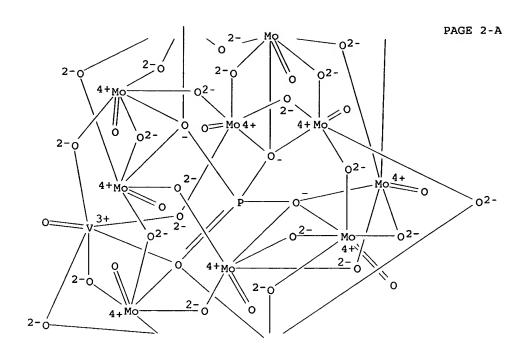
●6 Na+

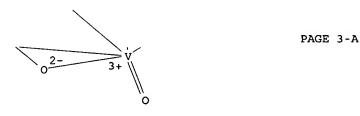
RN 122795-31-7 HCAPLUS
CN Vanadate(5-), (heptadeca-μ-oxodecaoxodecamolybdate)hepta-μoxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ
':κΟ':κΟ':κΟ'':κΟ'':κΟ'''
:κΟ''':κΟ''']]di-, pentapotassium (9CI) (CA INDEX
NAME)





●5 K+

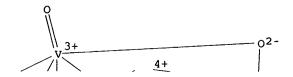


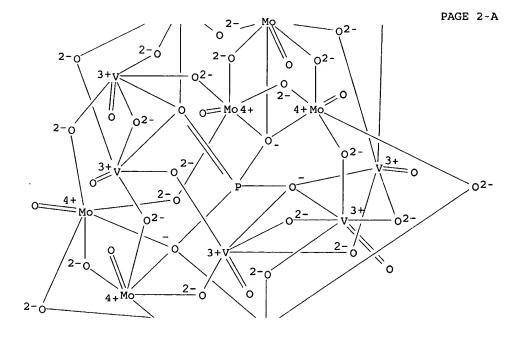


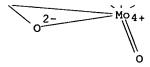
●5 Li+

RN 187289-60-7 HCAPLUS
CN Vanadate(9-), (octa-μ-oxohexaoxohexamolybdate)hexadeca-μoxohexaoxo[μ12-[phosphato(3-)-κΟ:κΟ:.kappa
.0':κΟ':κΟ':κΟ'':κΟ'':κΟ'
'':κΟ''':κΟ''']]hexa-, nonasodium (9CI) (CA INDEX
NAME)

PAGE 1-A



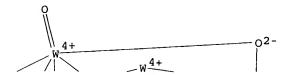




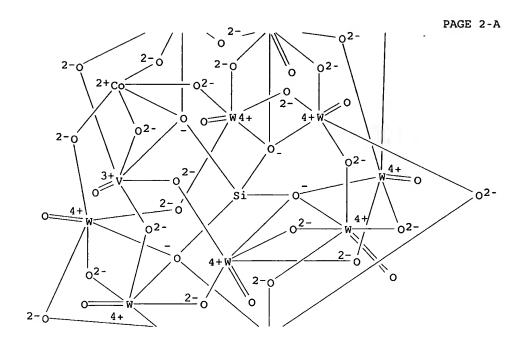
●9 Na+

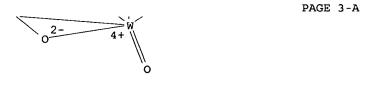
RN 273201-47-1 HCAPLUS
CN Vanadate(7-), cobaltate(heptadeca-μ oxodecaoxodecatungstate)[μ12-{orthosilicato(4-) κ0:κ0:κ0:κ0':κ0':κ0'
 ':κ0'':κ0'':κ0''':κ0''']hepta μ-οxooxo-, heptapotassium (9CI) (CA INDEX NAME)

PAGE 1-A



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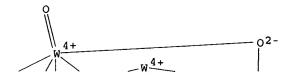


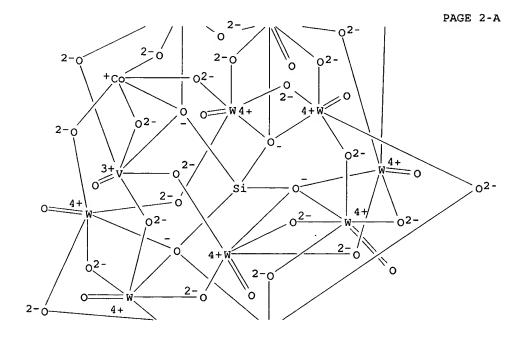


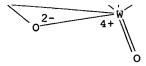
●7 K+

RN 340737-25-9 HCAPLUS
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 ':κΟ'':κΟ'':κΟ''':κΟ''']]heptaμ-oxooxo-, octapotassium (9CI) (CA INDEX NAME)

PAGE 1-A

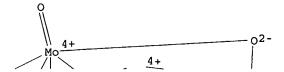


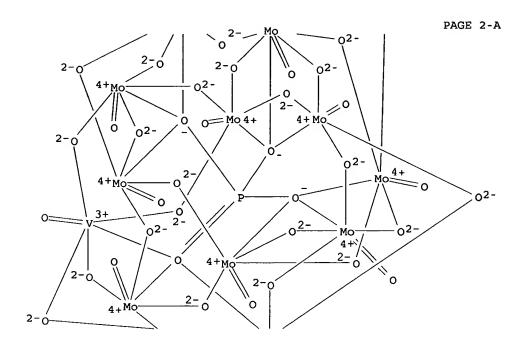


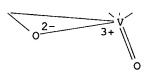


●8 K+

RN 340737-27-1 HCAPLUS
CN Vanadate(5-), (heptadeca-μ-oxodecaoxodecamolybdate)hepta-μoxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ
':κΟ':κΟ':κΟ'':κΟ'':κΟ'''
:κΟ''':κΟ''']]di-, pentasilver(1+) (9CI) (CA INDEX
NAME)

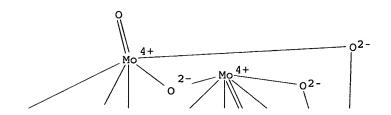


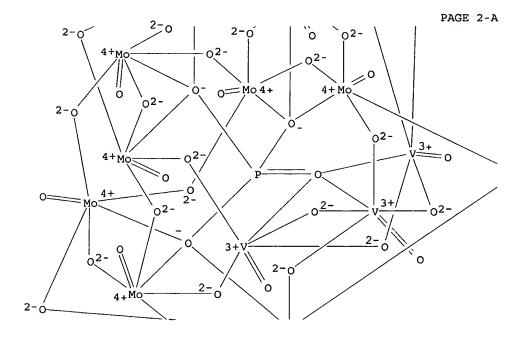




●5 Ag(I)+

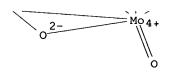
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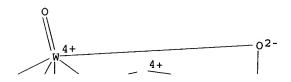


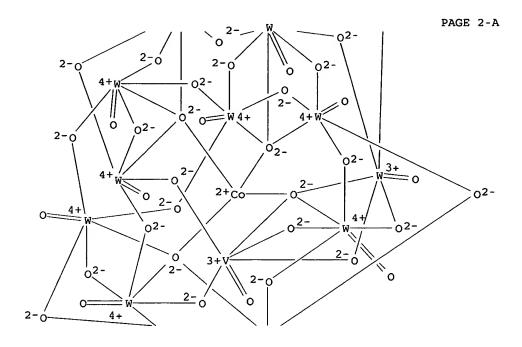
PAGE 3-A

●6 Ag(I)+

RN 340737-29-3 HCAPLUS

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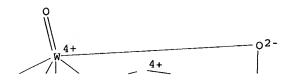


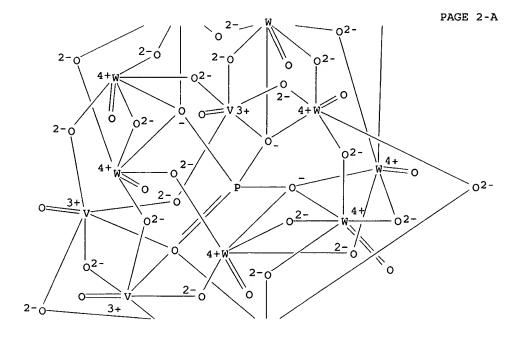


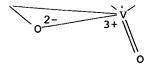
●8 Ag(I)+

RN 340737-30-6 HCAPLUS
CN Vanadate(7-), (dodeca-μ-oxooctaoxooctatungstate)dodeca-μoxotetraoxo[μ12-[phosphato(3-)-κ0:κ0:κ0:kapp
a.0':κ0':κ0':κ0'':κ0'':κ0'':κ0'
''':κ0''':κ0''']]tetra-, pentasodium dihydrogen (9CI)
(CA INDEX NAME)

PAGE 1-A



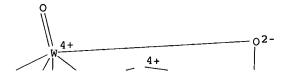


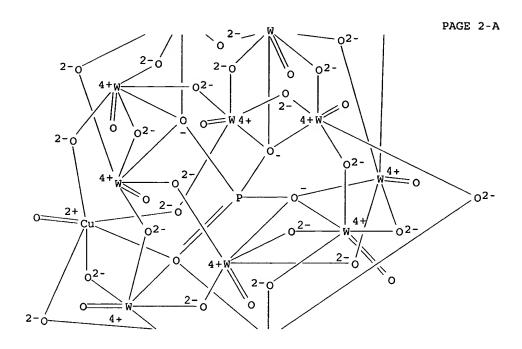


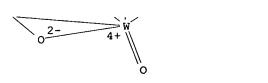
●2 H+

●5 Na+

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pa.O':κΟ'':κΟ'':κΟ'':.ka
ppa.O''']]undeca-, pentasodium (9CI) (CA INDEX NAME)



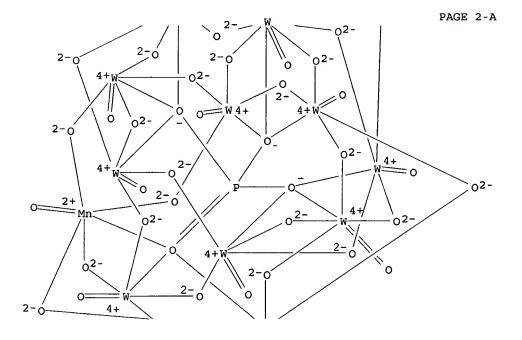


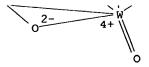


●5 Na+

RN 340737-33-9 HCAPLUS
CN Tungstate(5-), tetracosa-μ-oxoundecaoxo(oxomanganate) [μ12[phosphato(3-)-κΟ:κΟ:κΟ:κΟ':.kap
pa.O':κΟ'':κΟ'':κΟ''':.ka
ppa.O''']]undeca-, pentasodium (9CI) (CA INDEX NAME)







●5 Na+

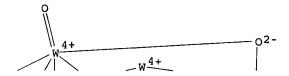
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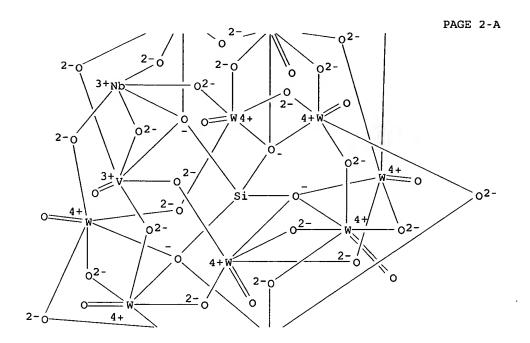
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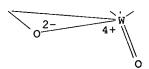
(polyoxometalate materials for removing environmental contaminant)

RN 340737-35-1 HCAPLUS

Ni 340/3/-35-1 RCAPBOS
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κΟ':κΟ'':κΟ'':κΟ'':κΟ''':κΟ'''
:κΟ'''] hepta-μ-oxo(oxovanadate)-, hexasodium (9CI) (CA INDEX NAME)







●6 Na+

IC ICM B01D053-00

CC 4-3 (Toxicology)

ST polyoxometalate environment contaminant

removal

IT Environmental pollution

Toxicants

(polyoxometalate materials for removing environmental contaminant)

IT Heteropoly acids

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)

(polyoxometalate materials for removing environmental contaminant)

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L114 ANSWER 34 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
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DOCUMENT NUMBER:
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                          agent detection and decontamination
AUTHOR (S):
                          Rhule, Jeffrey T.; Hill, Craig L.
CORPORATE SOURCE:
                          Department of Chemistry, Emory University,
                          Atlanta, GA, 30322, USA
SOURCE:
                          Proceedings of the ERDEC Scientific Conference
                          on Chemical and Biological Defense Research,
                          Aberdeen Proving Ground, MD, United States, Nov. 17-20, 1998 (1999), Meeting Date 1998,
                          307-313. Editor(s): Berg, Dorothy A.
                          National Technical Information Service:
                          Springfield, Va.
```

CODEN: 69AJH3 Conference English

The purpose of this research is to investigate the detection and decontamination capabilities of polyoxometalates

(POMs) when used in conjunction with currently available topical skin protectant creams (TSPs). H6[PV3Mo9O40] was finely ground and mixed in the cream to give a 25% weight/weight suspension.

CEES was layered on top of the cream and the time required for the POM-cream mixture to undergo a perceivable color change was noted. In new expts., the first systems for the rapid and catalytic oxidation degradation of CEES in a TSP model were discovered: POM/CEES/dichloroethane - trifluroethanol /oxidant, where the oxidant is O2 or benzoyl peroxide.

IT 12293-24-2

DOCUMENT TYPE:

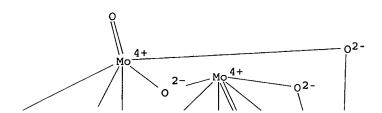
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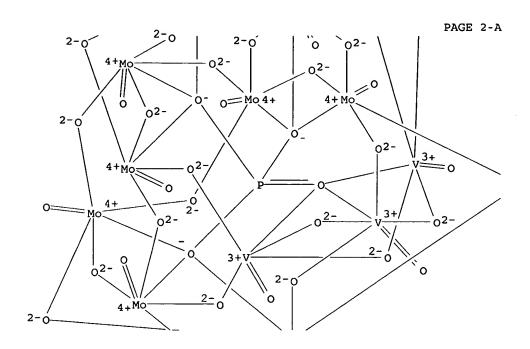
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(polyoxometalate topical skin protectant creams for chemical warfare agent detection and

decontamination)

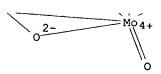
RN 12293-24-2 HCAPLUS





PAGE 2-B

)o2-



PAGE 3-A

●6 H+

CC 4-3 (Toxicology)

ST polyoxometalate chem warfare agent detection; decontamination chem warfare agent polyoxometalate; protection skin chem warfare

agent polyoxometalate
IT Chemical warfare agents

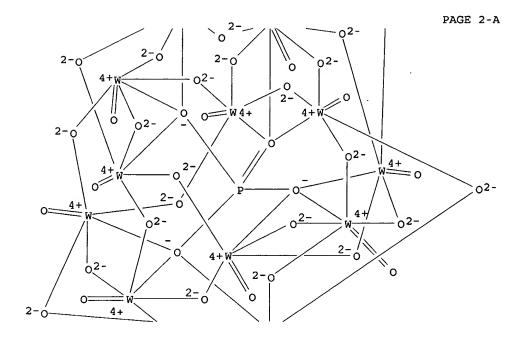
(polyoxometalate topical skin protectant creams for

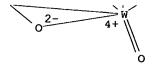
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chemical warfare agent detection and
        decontamination)
TΤ
     Heteropoly acids
     RL: BPR (Biological process); BSU (Biological study,
     unclassified); BUU (Biological use, unclassified); BIOL
     (Biological study); PROC (Process); USES (Uses)
         (polyoxometalate topical skin protectant creams for
        chemical warfare agent detection and
        decontamination)
ΙT
     Drug delivery systems
        (topical, PFPE #1511; polyoxometalate topical skin
        protectant creams for chemical warfare agent detection
        and decontamination)
     693-07-2, 2-Chloroethylethylsulfide
IT
     RL: ADV (Adverse effect, including toxicity); BIOL (Biological
        (polyoxometalate topical skin protectant creams for
        chemical warfare agent detection and
        decontamination)
TT
     12293-24-2
     RL: BAC (Biological activity or effector, except adverse); BSU
     (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (polyoxometalate topical skin protectant creams for
        chemical warfare agent detection and
        decontamination)
REFERENCE COUNT:
                                THERE ARE 9 CITED REFERENCES AVAILABLE
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L114 ANSWER 35 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                          2000:567386 HCAPLUS
DOCUMENT NUMBER:
                          133:210587
TITLE:
                          Preparation of high temperature
                          composite membranes for hydrogen
                         proton exchange membrane fuel cells
AUTHOR (S):
                          Lin, Jung-Chou; Kunz, H. Russell; Cutlip,
                         Michael B.; Fenton, James M.
CORPORATE SOURCE:
                         Department of Chemical Engineering, University
                         of Connecticut, Storrs, CT, 06269-3222, USA
SOURCE:
                         Hazardous and Industrial Wastes (1999), 31st,
                         656-662
                         CODEN: HIWAEB; ISSN: 1044-0631
PUBLISHER:
                         Technomic Publishing Co., Inc.
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     High temperature proton exchange membranes (PEM) for fuel cells have
     been prepared Single-cell testing results using methanol fuel
     verify that membranes can be operated at over 100°C without
     pressurizing the system. Internal resistance (IR) loss at
     120°C for Nafion-Teflon-phosphotungstic acid (NTPA)
     membrane is 19 mv at 108 mA/cm2 which is superior to other
     available membranes. Nafion-zirconium hydrogen phosphate (NZHP)
     membranes also show reasonable conductivity and favorable characteristics
     at temps. over 100 °C. Incorporating platinum into the NZP composite membrane is being conducted to further improve
     the conductivity Other potential high temperature membranes such as Nafion
     containing fine particles have been prepared for comparison. Higher
     temperature membranes should improve resistance to carbon monoxide
     poisoning.
TΤ
     1343-93-7, Phosphotungstic acid
     RL: DEV (Device component use); USES (Uses)
        (preparation of high temperature composite membranes for
        hydrogen proton exchange membrane fuel cells)
RN
     1343-93-7 HCAPLUS
CN
     Tungstate (3-), tetracosa-\mu-oxododecaoxo [\mu12-[phosphato(3-)-
```

κΟ:κΟ:κΟ':κΟ':κΟ':κΟ'
':κΟ'':κΟ'':κΟ''':κΟ''']]dodec
a-, trihydrogen (9CI) (CA INDEX NAME)

PAGE 1-A







●3 H+

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 38, 72 IT Fluoropolymers, uses Fluoropolymers, uses RL: DEV (Device component use); USES (Uses) (preparation of high temperature composite membranes for hydrogen proton exchange membrane fuel cells) ΤT Fuel cells (proton exchange membrane; preparation of high temperature composite membranes for hydrogen proton exchange membrane fuel cells) IT 1343-93-7, Phosphotungstic acid 9002-84-0, Teflon 66796-30-3, Nafion 117 RL: DEV (Device component use); USES (Uses)

(preparation of high temperature composite membranes for hydrogen proton exchange membrane fuel cells)

REFERENCE COUNT: 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE

IN THE RE FORMAT

L114 ANSWER 36 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:132682 HCAPLUS

DOCUMENT NUMBER:

133:1581

TITLE:

Polyoxometalate oxidation of

chemical warfare agent simulants in

fluorinated media

AUTHOR (S):

Johnson, Rhoma P.; Hill, Craig L.

CORPORATE SOURCE: Department of Chemistry, Emory University,

Atlanta, GA, 30322, USA

SOURCE: Journal of

Journal of Applied Toxicology (1999),

19(Suppl. 1), S71-S75

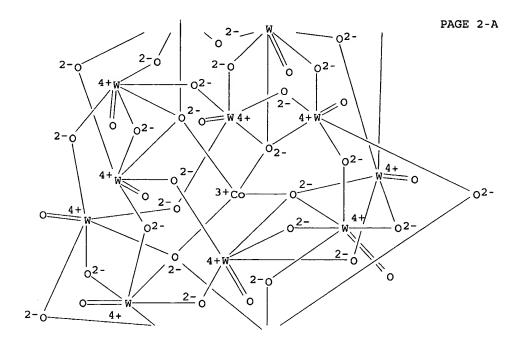
CODEN: JJATDK; ISSN: 0260-437X John Wiley & Sons Ltd.

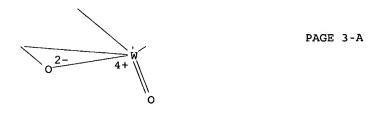
PUBLISHER: John Wiley & Sons

DOCUMENT TYPE: Journal LANGUAGE: English

The aim of this research is to determine if appropriate polyoxometalates (POMs) could be added to perfluoropolyether topical skin protectants (TSPs) currently available or under development to give these TSPs the addnl. capability of detecting and in some cases catalytically decontaminating sulfur mustard (HD) and perhaps other chemical warfare agents (CWAs) at ambient temps. Detection would be based on significant color changes in the POM upon reduction by the CWA whereas catalytic decontamination would be based on the ability of some families of POMs to catalyze O2-based oxidns. by more than one mechanism. Five POMs (10-25% by weight) were each suspended in .apprx.5 g of the perfluoropolyether (PFPE, CF30[-CF(CF3)CF20-]x(-CF20-)yCF3) "barrier" cream. A stoichiometric amount of HD sulfide simulant was layered on top of each POM-cream mixture The short reaction times were recorded for each system. Mechanistic studies were conducted using an PFPE oil analog of the barrier cream in a microemulsion with the sulfide simulant, POM, PFPE surfactant and

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

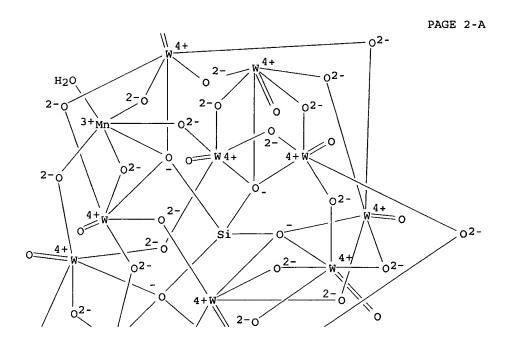




●5 K+

 $\cos a - \mu - oxoundecaoxoundeca-$, pentapotassium (9CI) (CA INDEX NAME)





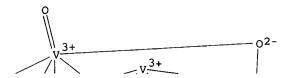
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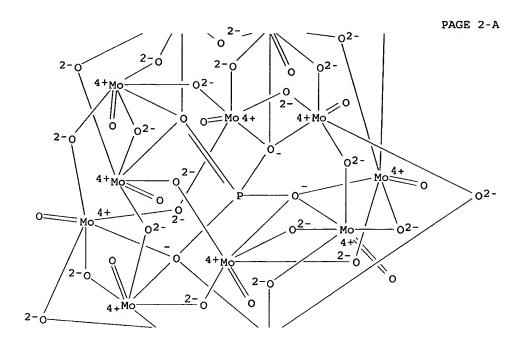
PAGE 3-A

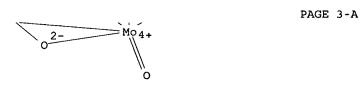
●5 K+

RN 222989-25-5 HCAPLUS
CN Vanadate(5-), (heptadeca-μ-oxodecaoxodecamolybdate)hepta-μoxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ
':κΟ':κ,Ο'κ:Ο''κ:Ο''κ:Ο''
'κ:Ο'''κ:Ο''']]di-, pentahydrogen (9CI) (CA INDEX

NAME)

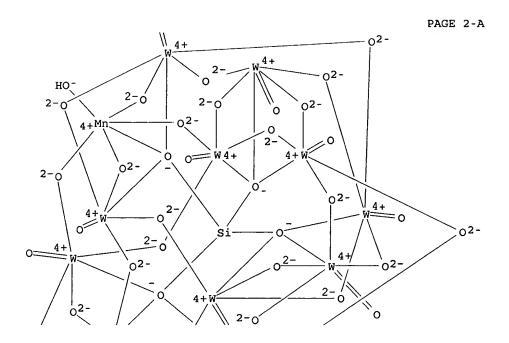


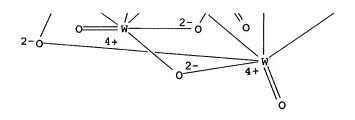




●5 H+







```
4-3 (Toxicology)
CC
ST
     polyoxometalate oxidn chem warfare agent
IT
     Polyethers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (fluorine-containing, topical skin protectants;
        polyoxometalate oxidation of chemical warfare agent
        simulants in fluorinated media)
TΤ
     Polyethers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (perfluoro, topical skin protectants; polyoxometalate
        oxidation of chemical warfare agent simulants in
        fluorinated media)
     Fluoropolymers, biological studies Fluoropolymers, biological studies
IT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (polyether-, topical skin protectants; polyoxometalate
        oxidation of chemical warfare agent simulants in
        fluorinated media)
IT
     Chemical warfare agents
        (polyoxometalate oxidation of chemical warfare
        agent simulants in fluorinated media)
ΙT
     Heteropoly acids
     RL: BAC (Biological activity or effector, except adverse); BSU
     (Biological study, unclassified); BIOL (Biological study)
        (polyoxometalate oxidation of chemical warfare
        agent simulants in fluorinated media)
     12520-46-6 141503-78-8 222989-25-5
TT
     270252-09-0
     RL: ARG (Analytical reagent use); BUU (Biological use,
     unclassified); PRP (Properties); ANST (Analytical study); BIOL
     (Biological study); USES (Uses)
        (polyoxometalate oxidation of chemical warfare
        agent simulants in fluorinated media)
ŤΤ
     505-60-2, Sulfur mustard
     RL: RCT (Reactant); REM (Removal or disposal); PROC (Process);
     RACT (Reactant or reagent)
        (polyoxometalate oxidation of chemical warfare
        agent simulants in fluorinated media)
REFERENCE COUNT:
                                THERE ARE 19 CITED REFERENCES AVAILABLE
                         19
                                FOR THIS RECORD. ALL CITATIONS AVAILABLE
                                IN THE RE FORMAT
L114 ANSWER 37 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1999:375636 HCAPLUS
DOCUMENT NUMBER:
                         131:20590
TITLE:
                         Polyoxometalate bleach catalysts in
                         cleaning and detergent
                         compositions
INVENTOR(S):
                         Greenhill-Hooper, Michael John; Rey-Garcia,
                         Fernando; Corma-Canos, Avelino; Jorda-Moret,
```

Jose Luis

PATENT ASSIGNEE(S): U.S. Borax Inc., USA
SOURCE: PIXXD2

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

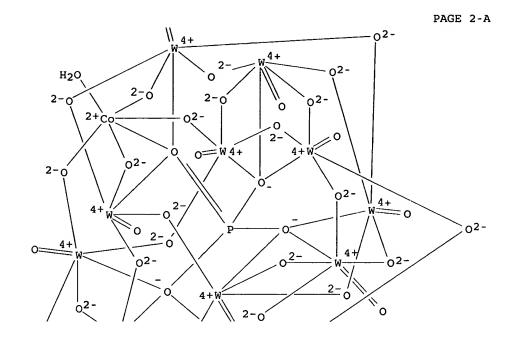
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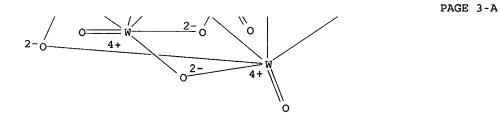
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WO 9928	- 426	A1	19990610	WO 1998-GB3618	1998
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ZA 9811				ZA 1998-11059	1998
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				AU 1999-13457	1998 1203
AU 9913	457	ΑI	19990616	AU 1999-13457	1998 1203
EP 1036	154	A1	20000920	EP 1998-957033	1998
R:	AT, BE, CH, MC, PT, IE,		ES, FR,	GB, GR, IT, LI, LU,	1203 NL, SE,
BR 9815		A	20001024	BR 1998-15343	1998
JP 2001	525451	T2	20011211	JP 2000-523306	1203
US 6326	342	B1	20011204	US 2000-555696	1998 1203
DD 100 1 MW 1 DD				an 1005 occ.	2000 0822
PRIORITY APP	LN. INFO.:			GB 1997-25614	A 1997 1203
				WO 1998-GB3618	W 1998 1203

AB A bleaching composition comprises (i) a bleaching agent such as peroxide, and (ii) bleach catalyst, a polyoxometalate of Keggin, Dawson or Finke structure (A')a'(Cox'Yy',Mm',Oo).cH2O, where A' = cation; a' has a value such that (A')a' counters the anionic charge of (Cox', Yy', Mm', Oo); x' = 0.25-4; Y = P, Si or Co; y' = 1 or 2; o = 34-68; M = W, Mo, V, Nb or Ta; m' = 9-18; and c = 0-84. The bleaching compns. according to the invention have good bleaching performance and can be used with or without a bleach activator, e.g. at low temps. Catalyst K6[Si(Co.H2O)W11039] was prepared

IT 39292-26-7P 105785-76-0P 226422-93-1P

```
226422-94-2P 226422-97-5P 226422-98-6P
    226422-99-7P 226423-00-3P
    RL: CAT (Catalyst use); IMF (Industrial manufacture); PREP
    (Preparation); USES (Uses)
       (bleach catalyst; bleaching compns. containing cobalt
       polyoxometalate bleach catalyst in laundering fabrics)
RN
    39292-26-7 HCAPLUS
    CN
    O''':κO''':κO''']]undecatungstate(5-) (5:1) (9CI) (CA
    INDEX NAME)
    CM
        1
    CRN 66258-00-2
    CMF Co H2 O40 P W11
CCI CCS
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CM 2

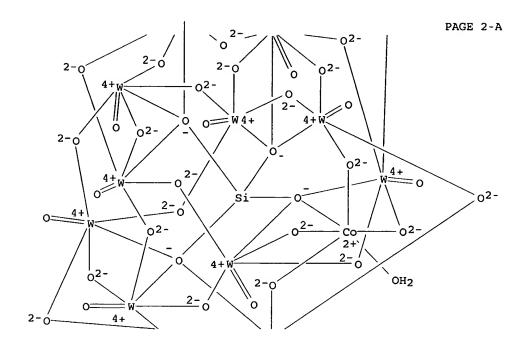
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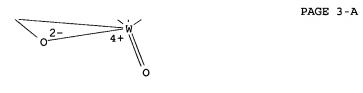
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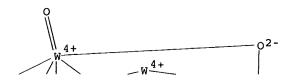
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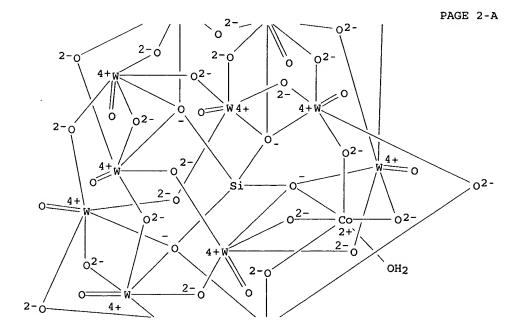
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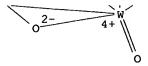




●6 K+



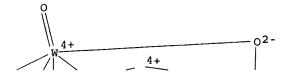


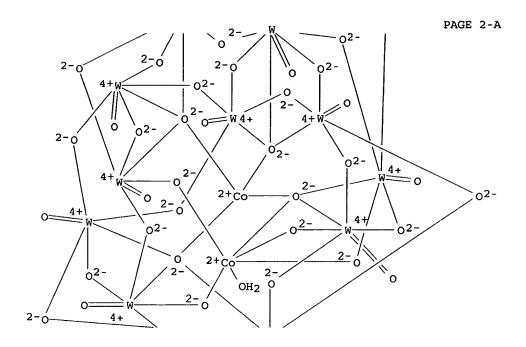


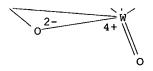
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RN CN

226422-94-2 HCAPLUS
Tungstate(8-), (aquadicobaltate)tetracosa-μ-oxotetra-μ4oxoundecaoxoundeca-, heptapotassium hydrogen (9CI) (CA INDEX
NAME)







● H+

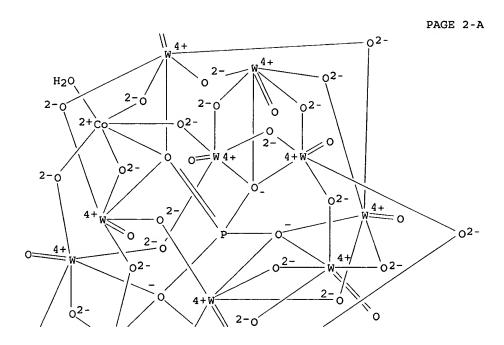
●7 K+

RN 226422-97-5 HCAPLUS
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pa.O':κΟ':κΟ':κΟ'':κΟ'':κ
O''':κΟ''':κΟ''']]undecatungstate(5-) (5:1) (9CI) (CA
INDEX NAME)

CM 1

CRN 66258-00-2
CMF CO H2 O40 P W11
CCI CCS





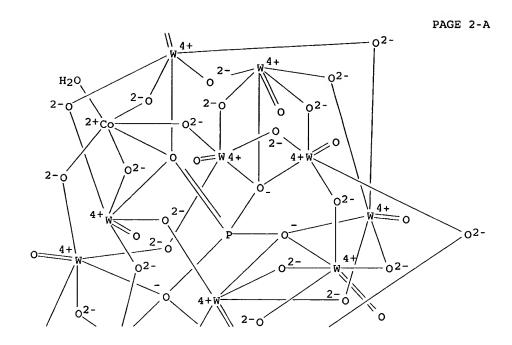
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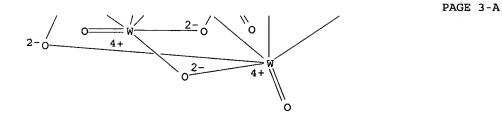
CRN 302-57-8 CMF C7 H18 N

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pa.O':κΟ':κΟ'':κΟ'':κΟ'':κ
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INDEX NAME)

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CRN 66258-00-2 CMF Co H2 O40 P W11 CCI CCS





CM 2

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RN 226422-99-7 HCAPLUS

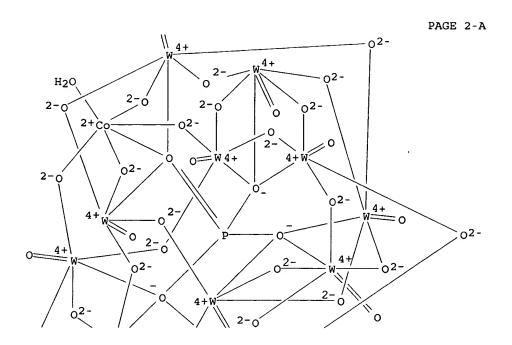
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INDEX NAME)

CM 1

CRN 66258-00-2 CMF CO H2 O40 P W11 CCI CCS

PAGE 1-A

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CM 2

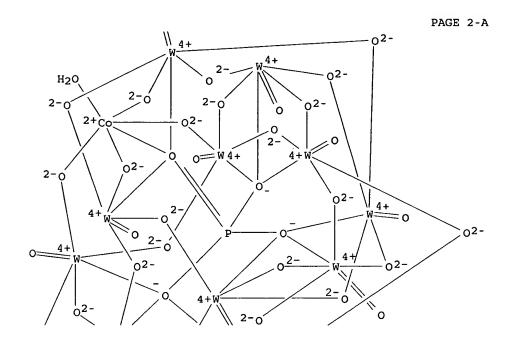
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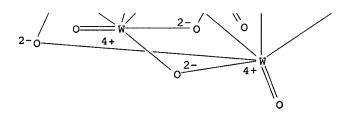
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CRN 66258-00-2 CMF Co H2 O40 P W11 CCI CCS

PAGE 1-A

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CM 2

CRN 10182-92-0 CMF C17 H38 N

 $Me_3+N-(CH_2)_{13}-Me$

ΙT 12027-38-2

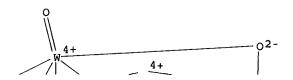
RL: RCT (Reactant); RACT (Reactant or reagent) (in catalyst preparation; bleaching compns. containing cobalt polyoxometalate bleach catalyst in laundering fabrics)
12027-38-2 HCAPLUS
Tungstate(4-), [µ12-[orthosilicato(4-)-

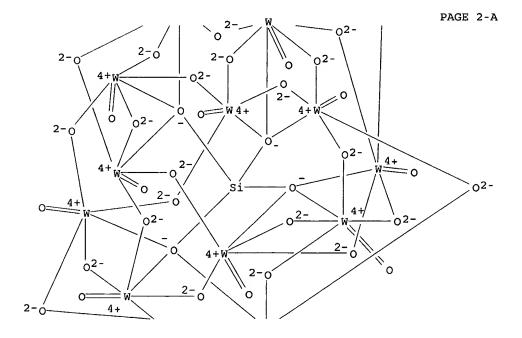
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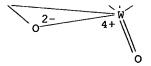
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':κ0'':κ0'':κ0''':κ0''':κ0''']]tetra

cosa-µ-oxododecaoxododeca-, tetrahydrogen (9CI) (CA INDEX NAME)







4 H+

```
TC
     ICM C11D003-39
     ICS C11D003-395
CC
     46-5 (Surface Active Agents and Detergents)
     Section cross-reference(s): 67
     cobalt polyoxometalate bleach catalyst; detergent bleach
ST
     polyoxometalate catalyst
IT
     Bleaching agents
     Catalysts
        (bleaching compns. containing cobalt
        polyoxometalate bleach catalyst in laundering fabrics)
IT
     Detergents
        (dishwashing; bleaching compns. containing cobalt
        polyoxometalate bleach catalyst in)
IT
     Detergents
        (laundry; bleaching compns. containing cobalt
        polyoxometalate bleach catalyst in laundering fabrics)
TΤ
     39292-26-7P 105785-76-0P 226422-93-1P
     226422-94-2P 226422-95-3P 226422-96-4P
     226422-97-5P 226422-98-6P 226422-99-7P
     226423-00-3P
     RL: CAT (Catalyst use); IMF (Industrial manufacture); PREP
     (Preparation); USES (Uses)
        (bleach catalyst; bleaching compns. containing cobalt
        polyoxometalate bleach catalyst in laundering fabrics)
IT
     3313-92-6, Sodium percarbonate 7722-84-1, Hydrogen peroxide,
     uses 11138-47-9, Sodium perborate
RL: TEM (Technical or engineered material use); USES (Uses)
        (bleach; bleaching compns. containing cobalt
        polyoxometalate bleach catalyst in laundering fabrics)
IT
     64-20-0, Tetramethylammonium bromide 71-48-7, Cobalt (II)
              127-08-2, Potassium acetate 298-14-6, Potassium
     acetate
     carbonate (KHCO3) 1643-19-2, Tetrabutylammonium bromide
     1941-30-6, Tetrapropylammonium bromide 2700-16-5,
     Triethylmethylammonium bromide 2840-24-6, Trimethylammonium
     bromide 12027-38-2 13472-45-2, Disodium tungstate
     81205-57-4
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (in catalyst preparation; bleaching compns. containing cobalt
        polyoxometalate bleach catalyst in laundering fabrics)
REFERENCE COUNT:
                               THERE ARE 2 CITED REFERENCES AVAILABLE
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                               FOR THIS RECORD. ALL CITATIONS AVAILABLE
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L114 ANSWER 38 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
                         1999:297315 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         130:332869
TITLE:
                         Polyoxometalate antifiloviral
                         composition containing a
                         heteropolytungstate, and preparation thereof
                         Matthews, Barry Ross; Holan, George
INVENTOR(S):
PATENT ASSIGNEE(S):
                         Starpharma Limited, Australia
                         PCT Int. Appl., 37 pp.
SOURCE:
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CODEN: PIXXD2

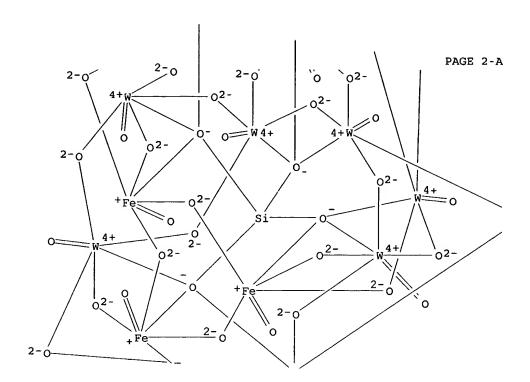
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                                                                                1024
                                                    WO 1998-AU880
                                                                                1998
                                                                                1023
     A method of prophylactic or therapeutic inhibition of Ebola virus
AB
     and other filoviruses in a human or non-human animal patient is
     provided which comprises administering to the patient an effective
     amount of a heteropolytungstate selected from one of the following
     formulas: AnM1-4WqOr, AnYMXW11039, An[(FeOA)4P2W18068],
     An[Co(OH)3(H2O)6(HPO4)2(P3W27O102)], or AnP2W15O56 (A = cation; n
      = number of cations for elec. neutrality; Y = ligand; X = B, P, Si,
     Ge, Zn, Co, Fe, Ga, Ti, Zr, V, Cu; M = various metals or
     combinations thereof; q=9-11, 15-18, 22, 30, 34; and r=39, 40, 56, 62, 65, 68, 78, 102, 112, 122). Preparation of heteropolytung state compds. is described.
ΙT
     152313-58-1
     RL: BAC (Biological activity or effector, except adverse); BSU
      (Biological study, unclassified); THU (Therapeutic use); BIOL
      (Biological study); USES (Uses)
         (polyoxometalate antifiloviral composition
         containing heteropolytungstate, and preparation thereof)
RN
     152313-58-1 HCAPLUS
     Tungstate(13-), [\mu 12-[orthosilicato(4-)-
CN
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      ':κ0'':κ0'':κ0''':κ0''':κ0''']]henei
     cosa-μ-oxononaoxo(tri-μ-oxotrioxotriferrate)nona-,
     hexapotassium heptahydrogen (9CI) (CA INDEX NAME)
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Patent

English

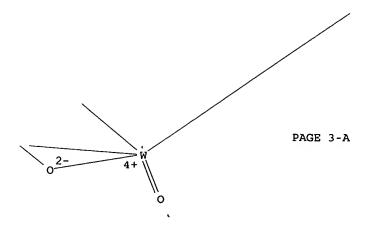
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LANGUAGE:



PAGE 2-B

)o2-



●7 H+

●6 K+

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     ICS
CC
     1-5 (Pharmacology)
     Section cross-reference(s): 63, 78
ST
    virucide filovirus heteropolytungstate prepn;
     polyoxometalate virucide filovirus; Ebola virus virucide
     heteropolytungstate prepn
TT
    Drug delivery systems
        (4polyoxometalate antifiloviral composition containing
        heteropolytungstate, and preparation thereof)
TT
     Fever and Hyperthermia
        (hemorrhagic, viral; polyoxometalate antifiloviral
        composition containing heteropolytungstate, and preparation thereof)
IT
    Antiviral agents
    Drug screening
    Ebola virus
     Filovirus
    Marburg virus
        (polyoxometalate antifiloviral composition
        containing heteropolytungstate, and preparation thereof)
IT
    Heteropoly acids
    RL: BAC (Biological activity or effector, except adverse); BSU
     (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (polyoxometalate antifiloviral composition
        containing heteropolytungstate, and preparation thereof)
TΤ
    Heteropoly acids
    RL: BAC (Biological activity or effector, except adverse); BSU
     (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (tungstates; polyoxometalate antifiloviral
        composition containing heteropolytungstate, and preparation thereof)
ΙT
    Drugs
        (veterinary; polyoxometalate antifiloviral
        composition containing heteropolytungstate, and preparation thereof)
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                  84750-84-5P
                               110717-64-1P 152444-40-1P
    179160-07-7P, Iron potassium tungsten oxide phosphate
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(Fe4K12W16O57(PO4)2)

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RL: BAC (Biological activity or effector, except adverse); BSU
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     (Therapeutic use); BIOL (Biological study); PREP (Preparation);
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        containing heteropolytungstate, and preparation thereof)
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         (preparation and reaction; polyoxometalate antifiloviral
        composition containing heteropolytungstate, and preparation thereof)
IT
     497-19-8, Carbonic acid disodium salt, reactions 7447-40-7,
     Potassium chloride, reactions 7601-89-0, Sodium perchlorate 7773-01-5, Manganese dichloride 10141-05-6, Cobalt dinitrate
     10421-48-4, Ferric nitrate 13138-45-9, Nickel dinitrate
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     RL: RCT (Reactant); RACT (Reactant or reagent)
         (reaction; polyoxometalate antifiloviral
        composition containing heteropolytungstate, and preparation thereof)
                                 THERE ARE 6 CITED REFERENCES AVAILABLE
REFERENCE COUNT:
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L114 ANSWER 39 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
                          1998:684433 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                          129:339856
TITLE:
                          Method, compositions, and aerosol
                           spray containing a polyoxometalate
                           for treating and preventing respiratory viral
                           infections
INVENTOR(S):
                           Schinazi, Raymond F.; Hill, Craig L.
PATENT ASSIGNEE(S):
SOURCE:
                           U.S., 18 pp., Cont.-in-part of U.S. Ser. No.
                           312,561, abandoned.
                           CODEN: USXXAM
DOCUMENT TYPE:
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LANGUAGE:
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FAMILY ACC. NUM. COUNT:
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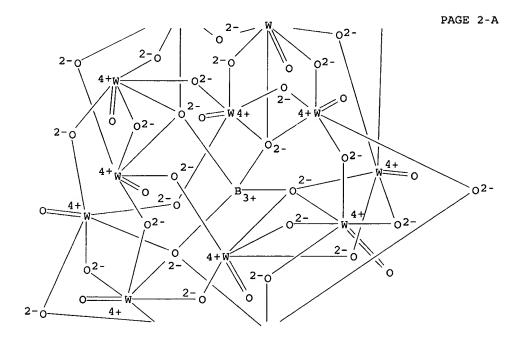
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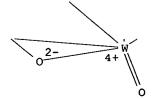
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     Respiratory viral infections may be effectively prevented or
     treated by administering an aerosol spray comprising a
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     infected primary human PBM cells and had no cytotoxicity to
     uninfected human PBM cells when evaluated up to 100 \mu\text{M}.
     11078-54-9 12027-38-2D, solid solution with
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     with tetrahydrogen analog 81552-97-8 82679-05-8
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     101144-77-8 112763-08-3 112763-08-3D,
     solid solution with tetrahydrogen analog 119390-04-4
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     139631-93-9 139631-95-1 139631-96-2
     139631-98-4 139632-00-1 141483-63-8
     162958-09-0 162958-11-4 162958-14-7
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     RL: BAC (Biological activity or effector, except adverse); BSU
     (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (method and aerosol spray containing a polyoxometalate
        for treating and preventing respiratory viral infections)
RN
     11078-54-9 HCAPLUS
CN
     Tungstate(5-), tetracosa-μ-oxododecaoxo[μ12-
     [tetrahydroxyborato(5-)-κ0:κ0:κ0:κ0':.kapp
     a.0':κ0':κ0'':κ0'':κ0'':κ0''':.kappa
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.O''':κO''']]dodeca-, pentapotassium (9CI) (CA INDEX NAME)

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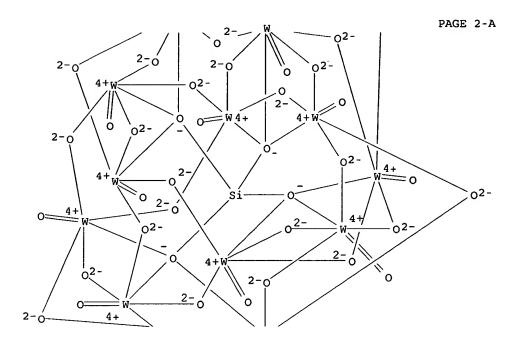


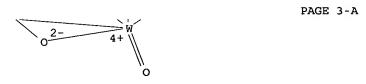




●5 K+

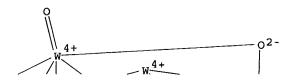


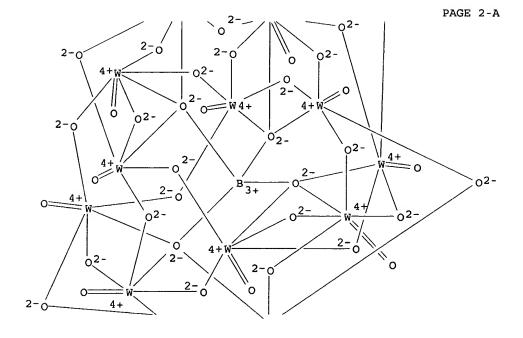


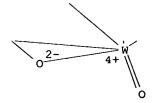


●4 H+

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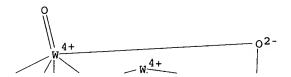


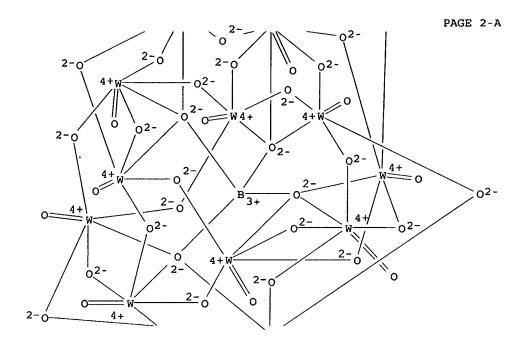




●5 Na+

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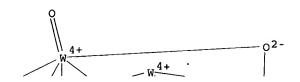


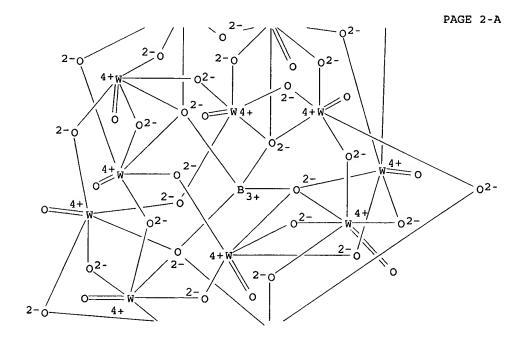


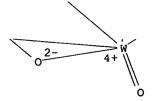


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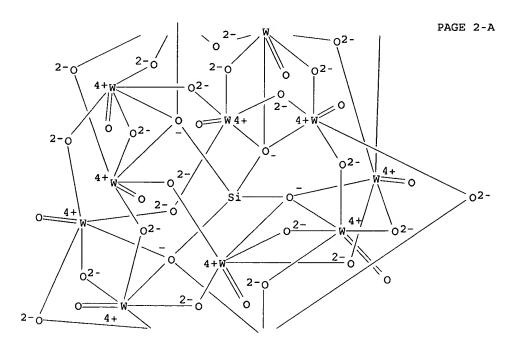
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CN Tungstate(5-), tetracosa-μ-oxododecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ::.kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':.kappa
.O''':κΟ''']]dodeca-, pentahydrogen (9CI) (CA INDEX NAME)

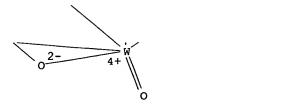




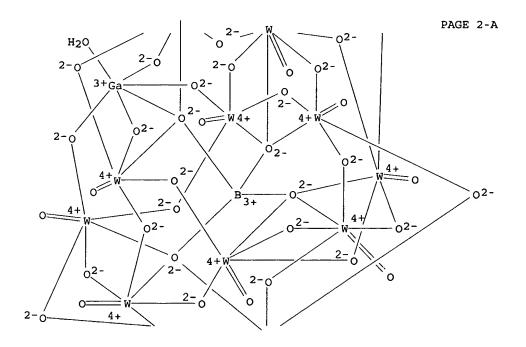


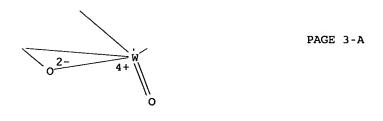
●5 H+





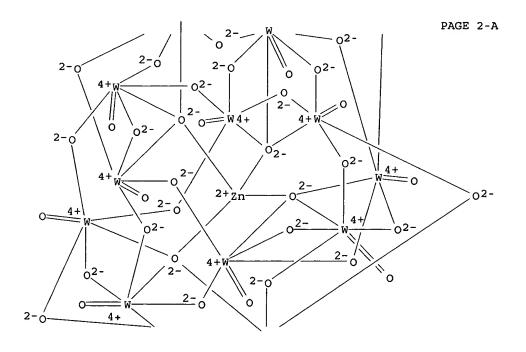
●4 NH4+

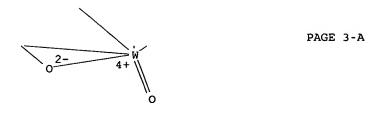




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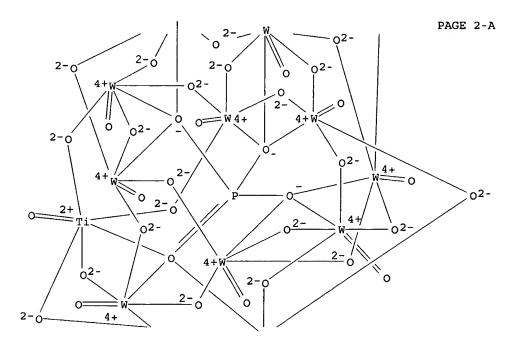
RN 82679-05-8 HCAPLUS CN Tungstate(6-), tetracosa- μ -oxotetra- μ 4-oxododecaoxozincatedodeca-, hexahydrogen (9CI) (CA INDEX NAME)

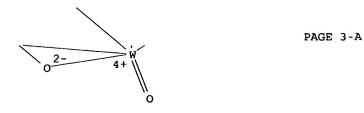




●6 H+

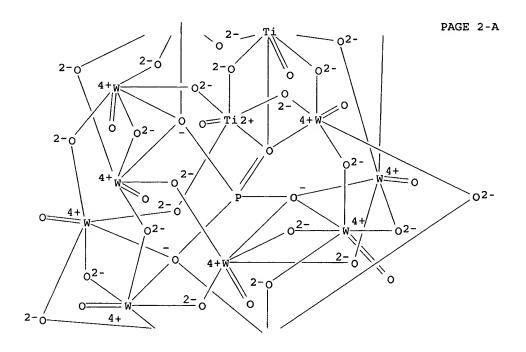
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CN Titanate(5-), (eicosa-μ-oxoundecaoxoundecatungstate)tetra-μoxooxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ':
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kappa.O''':κΟ''']]-, pentacesium (9CI) (CA INDEX NAME)

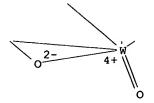




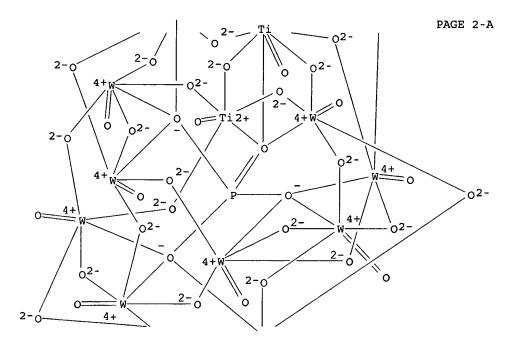
●5 Cs+

RN 84303-03-7 HCAPLUS
CN Titanate(7-), (heptadeca-μ-oxodecaoxodecatungstate)hepta-μoxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ
':κΟ':κΟ':κΟ'':κΟ'':κΟ'''
:κΟ''':κΟ''']}di-, heptacesium (9CI) (CA INDEX NAME)





●7 Cs+



CM 2 CRN 51-92-3

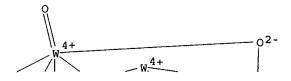
CMF C4 H12 N

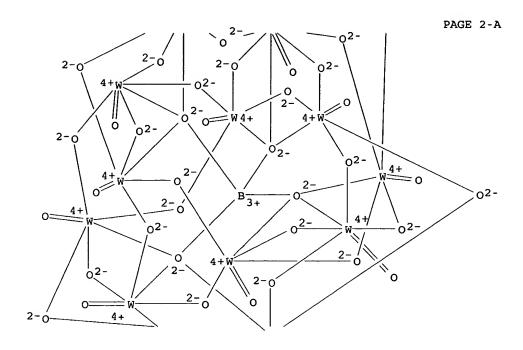
101144-77-8 HCAPLUS RN $Vanadate \, (9\text{-}) \, , \quad [\text{heptacosa-}\mu\text{-}oxopentadecaoxo} \, [\mu 9\text{-}[\text{phosphato} \, (3\text{-})\text{-}] \,]$ CN κ0:κ0:κ0:κ0':κ0':κ0'',κ0 $'':\kappa0''':\kappa0''']$]pentadecatungstate]nona- μ oxotrioxo[μ9-[phosphato(3-)-κ0:κ0:κ0:κ0 ':κ0':κ0'':κ0'':κ0''':κ0''']]tri-, octapotassium hydrogen (9CI) (CA INDEX NAME)

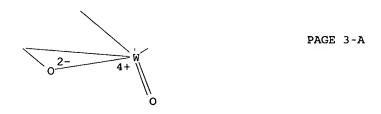
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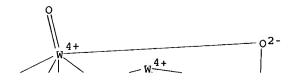


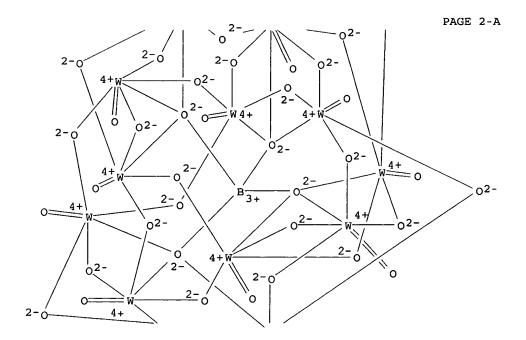


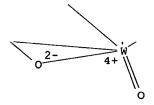
●5 NH₄+

RN 112763-08-3 HCAPLUS CN Tungstate(5-), tetracosa- μ -oxododecaoxo[μ 12-[tetrahydroxyborato(5-)- κ 0: κ 0: κ 0: κ 0':.kapp a.0': κ 0'': κ 0''': κ 0'': κ 0''': κ 0'': κ 0''': κ 0'''

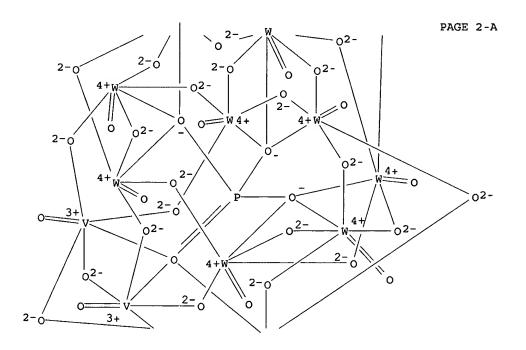
PAGE 1-A

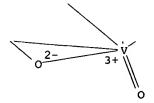






●5 NH4+



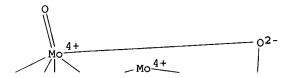


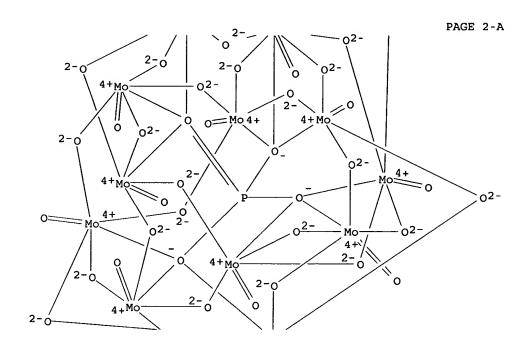
●6 K1

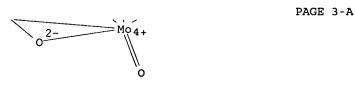
RN 129572-46-9 HCAPLUS
CN Molybdate(3-), tetracosa-μ-oxododecaoxo[μ12-[phosphato(3-)-κ0:κ0:κ0:κ0':κ0':κ0'':κ0'']]dodec
a-, trihydrogen, compd. with N,N-dimethylacetamide (1:3) (9CI) (CA INDEX NAME)

CM 1

CRN 12026-57-2
CMF H . 1/3 Mol2 O40 P
CCI CCS







●3 H+

CM 2

CRN 127-19-5 CMF C4 H9 N O

(9CI) (CA INDEX NAME)

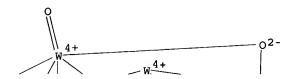
RN 131359-48-3 HCAPLUS
CN Tungstate(5-), tetracosa-μ-oxododecaoxo[μ12 [tetrahydroxyborato(5-)-κ0:κ0:κ0':.kapp
 a.0':κ0':κ0'':κ0'':κ0'':.kappa
 .0''':κ0''']]dodeca-, pentahydrogen (9CI) (CA INDEX NAME)

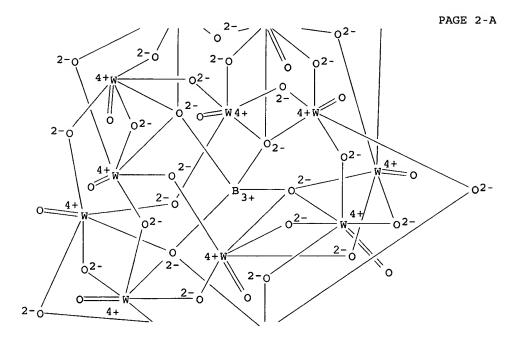
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN 131541-68-9 HCAPLUS
CN L-Lysine, tetracosa-μ-oxododecaoxo[μ12-[tetrahydroxyborato(5-)-0:0:0:0':0':0':0'':0'':0''':0''']]dodecatungstate(5-)

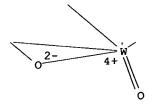
CM 1

CRN 12297-12-0 CMF B O40 W12 . 5 H

CCI CCS







●5 H+

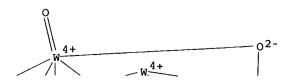
CM 2

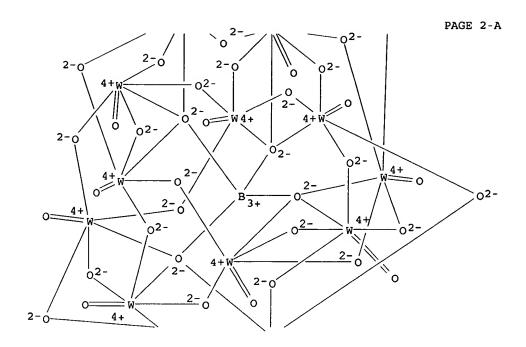
CRN 56-87-1 CMF C6 H14 N2 O2

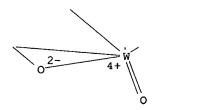
Absolute stereochemistry.

RN 131541-69-0 HCAPLUS
CN L-Arginine, tetracosa-μ-oxododecaoxo[μ12 [tetrahydroxyborato(5-)-κ0:κ0:κ0:κ0':.kapp
 a.0':κ0':κ0'':κ0'':κ0''':.kappa
 .0''':κ0''']]dodecatungstate(5-) (9CI) (CA INDEX NAME)
CM 1

CRN 12297-12-0 CMF B O40 W12 . 5 H CCI CCS







●5 H+

CM 2

CRN 74-79-3

CMF C6 H14 N4 O2

Absolute stereochemistry.

RN 131541-70-3 HCAPLUS

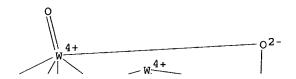
CN L-Histidine, tetracosa-µ-oxododecaoxo[µ12-

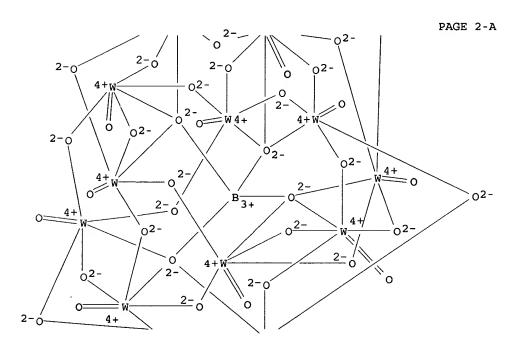
[tetrahydroxyborato(5-)-0:0:0:0':0':0':0'':0'':0'':0''':0''']
]dodecatungstate(5-) (9CI) (CA INDEX NAME)

CM 1

CRN 12297-12-0 CMF B O40 W12 . 5 H

CCI CCS





●5 H+

CM 2

CRN 71-00-1 CMF C6 H9 N3 O2

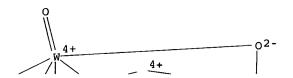
Absolute stereochemistry. Rotation (-).

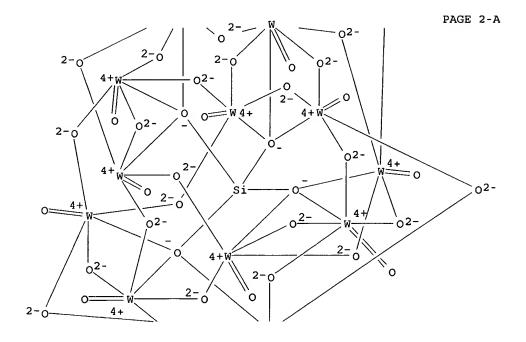
RN 132460-56-1 HCAPLUS

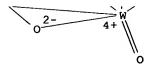
CN (CA INDEX NAME)

CM 1

CRN 12027-38-2 CMF H . 1/4 O40 Si W12 CCI CCS







●4 H+

CM 2

CRN 56-87-1 CMF C6 H14 N2 O2

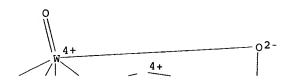
Absolute stereochemistry.

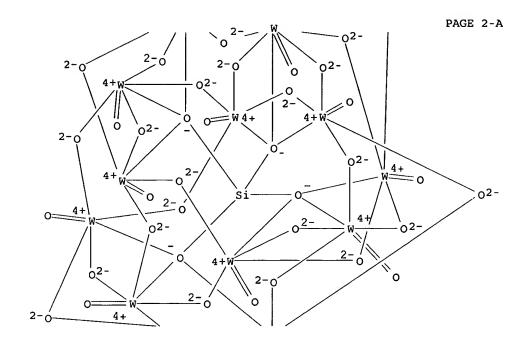
CM 1

CRN 12027-38-2

CMF H . 1/4 O40 Si W12

CCI CCS







●4 H+

CM 2

CRN 74-79-3 CMF C6 H14 N4 O2

Absolute stereochemistry.

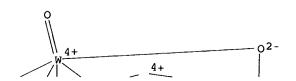
RN 132460-58-3 HCAPLUS
CN L-Histidine, [μ12-[orthosilicato(4-)κΟ:κΟ:κΟ:κΟ':κΟ':κΟ'
':κΟ'':κΟ'':κΟ''':κΟ''']]tetra
cosa-μ-oxododecaoxododecatungstate(4-) (9CI) (CA INDEX NAME)

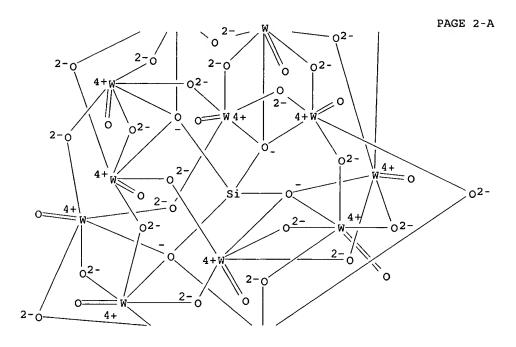
CM 1

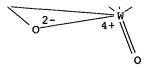
CRN 12027-38-2

CMF H . 1/4 040 Si W12

CCI CCS







■A 11+

CM 2

CRN 71-00-1 CMF C6 H9 N3 O2

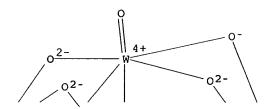
Absolute stereochemistry. Rotation (-).

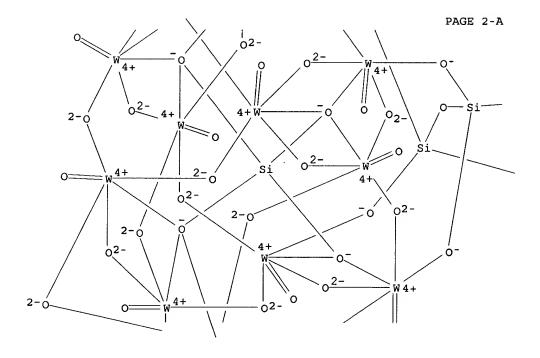
CMF C4 H6 O40 Si3 W11 CCI CCS

RN 138026-47-8 HCAPLUS
CN Methanaminium, N,N,N-trimethyl-, [μ4-[1,3-diethenyl-1,1,3,3-disiloxanetetrolato(4-)-κO1:κO1:κO3:κO3][
μ11-[orthosilicato(4-)-κO:κO:κO:κO':.ka
ppa.O':κO':κO'':κO'':κO'':κO''':.kap
pa.O''']]eicosa-μ-oxoundecaoxoundecatungstate(4-) (4:1) (9CI)
(CA INDEX NAME)

CM 1
CRN 137880-55-8

PAGE 1-A





* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

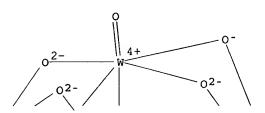
2-0 W 4+ 2-0 O O O

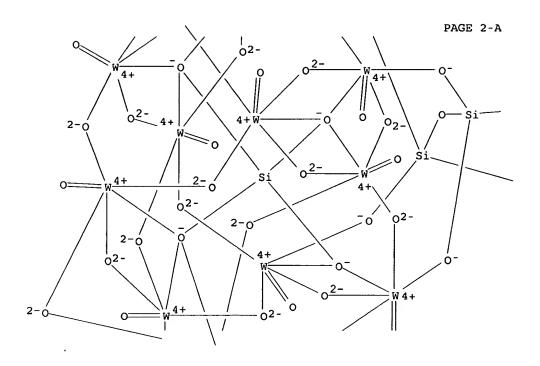
CM 2

CRN 51-92-3 CMF C4 H12 N

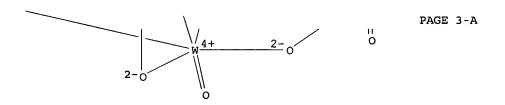
$$\begin{array}{c} \text{CH}_{3} \\ | \\ | \\ + \\ \text{CH}_{3} \\ \text{CH}_{3} \end{array}$$

RN 139631-93-9 HCAPLUS
CN Tungstate(4-), [μ11-[orthosilicato(4-)κΟ:κΟ:κΟ:κΟ':κΟ':κΟ'
':κΟ'':κΟ'':κΟ'''] eicosa-μoxoundecaoxo[μ4-[[4,4'-[1,1,3,3-tetra(hydroxy-κΟ)-1,3disiloxanediyl]bis[butanenitrilato]](4-)]]undeca-, tetracesium
(9CI) (CA INDEX NAME)





* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT



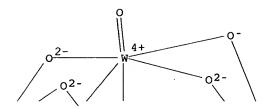
●4 Cs+

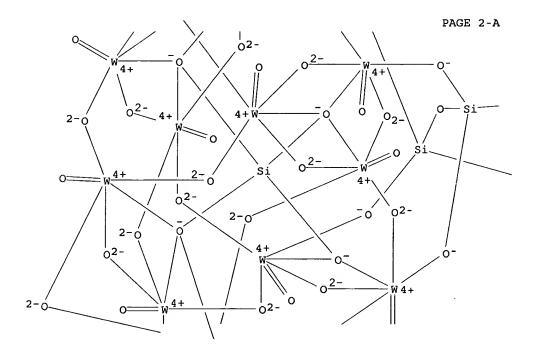
RN 139631-95-1 HCAPLUS
CN Methanaminium, N,N,N-trimethyl-, [μ11-[orthosilicato(4-)-κΟ:κΟ:κΟ:κΟ':κΟ':κΟ':κΟ'
':κΟ':κΟ':κΟ':κΟ':η]]eicosa-μoxoundecaoxo[μ4-[[4,4'-[1,1,3,3-tetra(hydroxy-κΟ)-1,3-disiloxanediyl]bis[butanenitrilato]](4-)]]undecatungstate(4-)(4:1) (9CI) (CA INDEX NAME)

CM 1

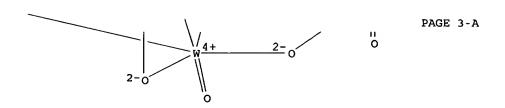
CRN 139631-94-0
CMF C8 H12 N2 O40 Si3 W11
CCI CCS

PAGE 1-A





* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

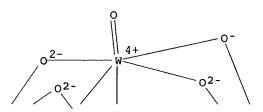


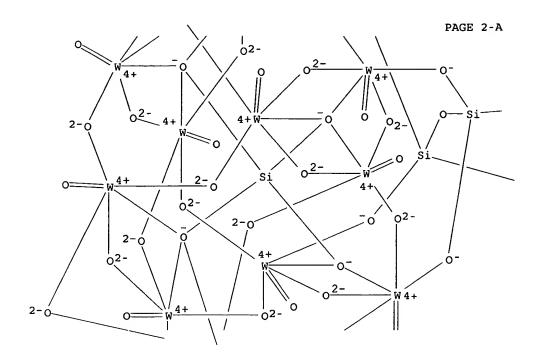
CM 2

CRN 51-92-3 CMF C4 H12 N

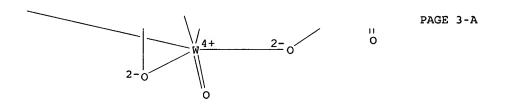
$$\begin{array}{c} \text{CH}_{3} \\ | \\ | \\ \text{H}_{3}\text{C} - \text{N} \xrightarrow{+} \text{CH}_{3} \\ | \\ \text{CH}_{3} \end{array}$$

RN 139631-96-2 HCAPLUS
CN Tungstate(4-), [μ4-[1,3-diethenyl-1,1,3,3-disiloxanetetrolato(4-)-κ01:κ01':κ03:κ03']] [μ11 [orthosilicato(4-)-κ0:κ0:κ0':κ0':κ0':κ0':κ0'':κ0'':κ0''': κ0''':κ0''':κ0'''
]]eicosa-μ-oxoundecaoxoundeca-, tetracesium (9CI) (CA INDEX NAME)



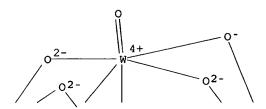


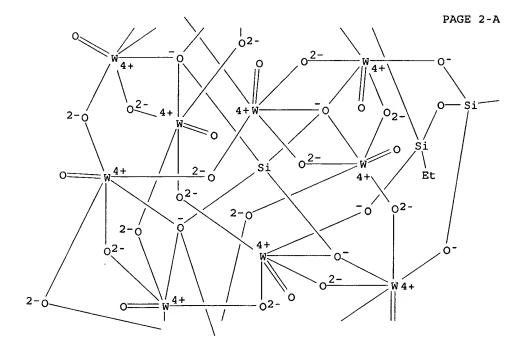
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT



●4 Cs+

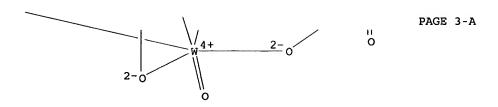
RN 139631-98-4 HCAPLUS
CN Methanaminium, N,N,N-trimethyl-, [μ4-[1,3-diethyl-1,1,3,3-disiloxanetetrolato(4-)-κ01:κ01':κ03:κ03']
] [μ11-[orthosilicato(4-)-κ0:κ0:κ0:κ0':.
kappa.0':κ0':κ0'':κ0'':κ0'':κ' κ0'':κ' κ0''':κ' κ0''':κ' κ0''':κ' κ0''':κ' κ0'''' κ0'''' κ0'''' κ0''' κ0'''' κ0''' κ0'''' κ0''' κ0''' κ0''' κ0''' κ0''' κ0''' κ0''' κ0''' κ0''' κ0'''' κ0''' κ0'''





PAGE 2-B

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CM 2

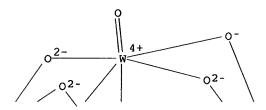
CRN 51-92-3 CMF C4 H12 N

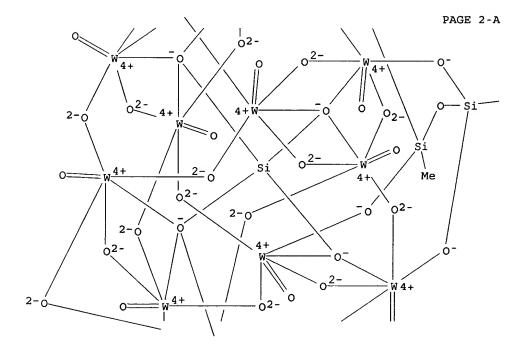
$$^{\text{CH}_3}_{\begin{array}{c} | \\ + \\ | \\ \text{CH}_3 \end{array}}$$

CM 1

CRN 139631-99-5 CMF C2 H6 O40 Si3 W11 CCI CCS

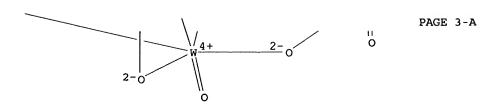
PAGE 1-A





PAGE 2-B

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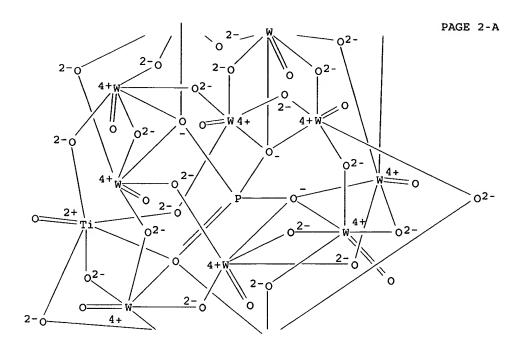
CM 2

CRN 51-92-3 CMF C4 H12 N

$$^{\text{CH}_3}_{|_{\frac{1}{N}}\text{CH}_3}$$

RN 141483-63-8 HCAPLUS
CN Titanate(5-), (eicosa-μ-oxoundecaoxoundecatungstate)tetra-μoxooxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ':
κΟ':κΟ'':κΟ'':κΟ''':
kappa.O''':κΟ''']]-, pentapotassium (9CI) (CA INDEX NAME)

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT



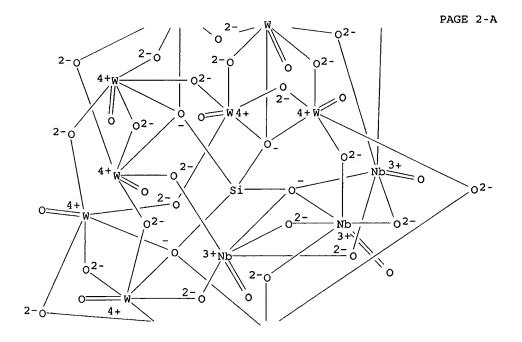


●5 K+

CCI CCS

PAGE 1-A





●7 H+

CM

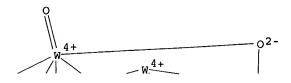
CRN 75-50-3 CMF C3 H9 N

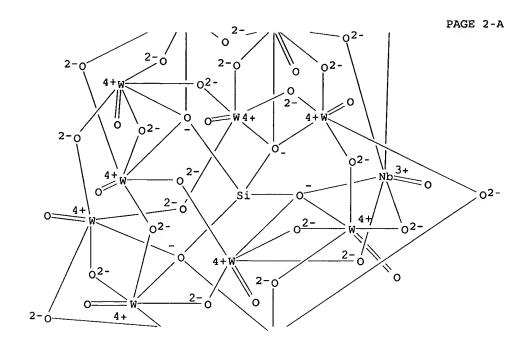
CH3 H3C-N-CH3

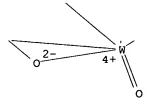
RN 162958-11-4 HCAPLUS Niobate(5-), (eicosa- μ -oxoundecaoxoundecatungstate)[μ 12-CN [orthosilicato(4-)-O:O:O':O':O':O':O'':O'':O''':O''']]tetr a-μ-οχοοχο-, pentahydrogen, compd. with N,N-dimethylmethanamine (1:5) (9CI) (CA INDEX NAME)

CM 1

CRN 162958-10-3 CMF H . 1/5 Nb O40 Si W11 CCI CCS







●5 H+

CM 2

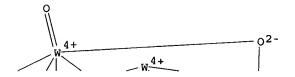
CRN 75-50-3 CMF C3 H9 N

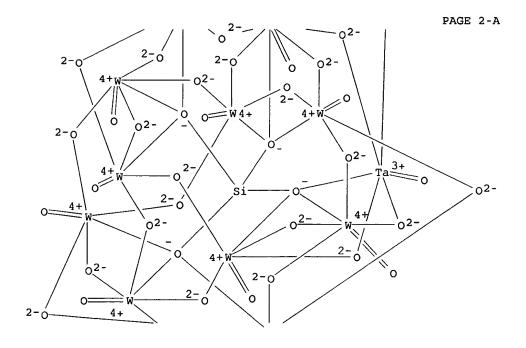
RN 162958-14-7 HCAPLUS

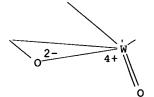
CN Tantalate(5-), (eicosa-μ-oxoundecaoxoundecatungstate) [μ12[orthosilicato(4-)-0:0:0':0':0':0'':0'':0'':0''']]tetr
a-μ-oxooxo-, pentahydrogen, compd. with N,N-dimethylmethanamine
(1:5) (9CI) (CA INDEX NAME)

CM 1

CRN 162958-13-6 CMF H . 1/5 O40 Si Ta W11 CCI CCS







●5 H+

CM 2

CRN 75-50-3 CMF C3 H9 N

CH₃ | H₃C-N-CH₃

RN 162958-20-5 HCAPLUS
CN Methanaminium, N,N,N-trimethyl-, [μ4-[1,3-dihexyl-1,1,3,3-disiloxanetetrolato(4-)-κΟ1:κΟ1:κΟ3:κΟ3]][
μ11-[orthosilicato(4-)-κΟ:κΟ:κΟ:κΟ':.ka
ppa.O':κΟ':κΟ'':κΟ'':κΟ'':kap
pa.O''']]eicosa-μ-oxoundecaoxoundecatungstate(4-) (4:1) (9CI)
(CA INDEX NAME)

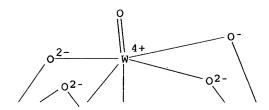
CM 1

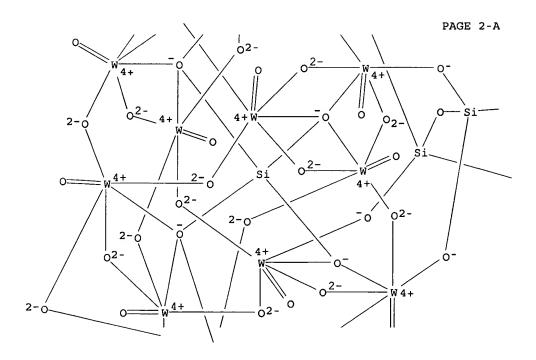
CRN 162958-19-2

CMF C12 H26 O40 Si3 W11

CCI CCS

PAGE 1-A





* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

CM 2

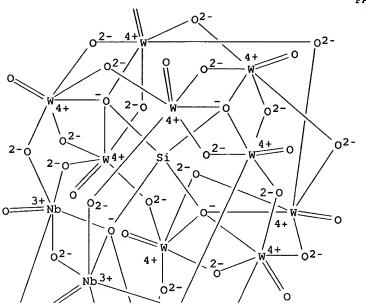
CRN 51-92-3 CMF C4 H12 N

$$\begin{array}{c} \text{CH3} \\ | \\ | \\ \text{H}_{3}\text{C--} \\ | \\ \text{CH}_{3} \end{array}$$

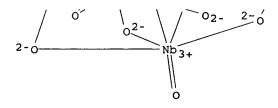
PAGE 1-A

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PAGE 2-A

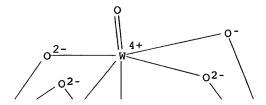


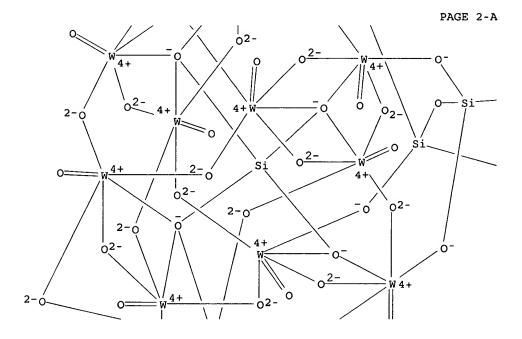
PAGE 3-A



●7 K+

RN 189823-27-6 HCAPLUS
CN Tungstate(4-), [μ4-[1,3-bis(3-chloropropyl)-1,1,3,3-disiloxanetetrolato(4-)-κ01:κ01':κ03:κ03']
] [μ11-[orthosilicato(4-)-κ0:κ0':κ0':.kappa.0':κ0':κ0'':κ0'':κ0'':.k
appa.0''']]eicosa-μ-oxoundecaoxoundeca-, tetracesium (9CI) (CA INDEX NAME)

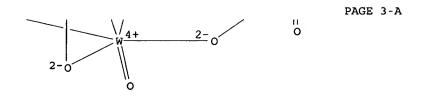




PAGE 2-B

$$---- (CH2)3-Cl$$

 \sim (CH₂)₃-Cl



●4 Cs+

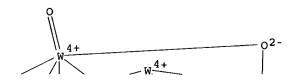
```
194150-76-0 HCAPLUS
RN
CN
     Tungstate(3-), tetracosa-\mu-oxododecaoxo[\mu12-[phosphato(3-)-
      κ0:κ0:κ0:κ0':κ0':κ0':κ0'
      ':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec
a-, trihydrogen, compd. with 1-methyl-2-pyrrolidinone (1:3) (9CI)
      (CA INDEX NAME)
```

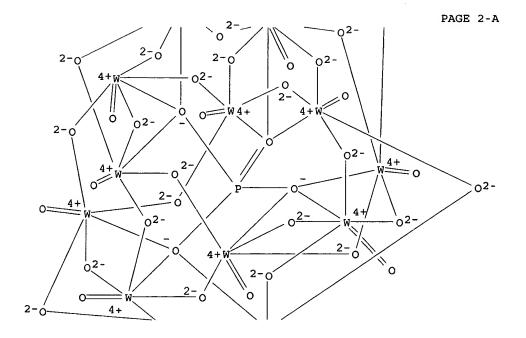
CM 1

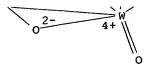
CRN 1343-93-7 CMF H . 1/3 O40 P W12 CCI CCS

Les Henderson

PAGE 1-A







●3 H+

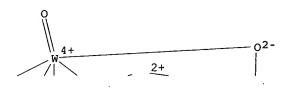
CM 2

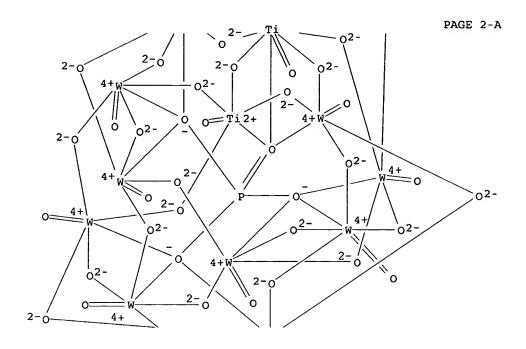
CRN 872-50-4 CMF C5 H9 N O

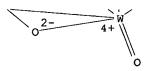


215594-65-3 HCAPLUS RNCN

Titanate(7-), (heptadeca- μ -oxodecaoxodecatungstate)hepta- μ -oxodioxo[μ 12-[phosphato(3-)- κ 0: κ 0: κ 0': κ 0







●7 Na+

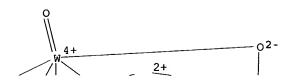
RN 215594-66-4 HCAPLUS
CN L-Arginine, (heptadeca-μ-oxodecaoxodecatungstate)hepta-μoxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ
':κΟ':κΟ':κΟ':κΟ'':κΟ'''
:κΟ''':κΟ''']]dititanate(7-) (7:1) (9CI) (CA INDEX
NAME)

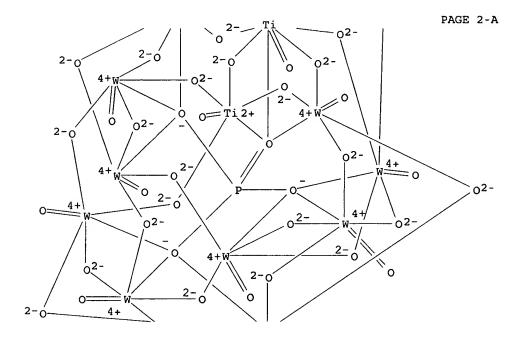
CM 1

CRN 215601-31-3 CMF H . 1/7 O40 P Ti2 W10

CCI CCS

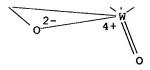
PAGE 1-A





571-272-2538

PAGE 3-A



●7 H⁺

CM 2

CRN 74-79-3 CMF C6 H14 N4 O2

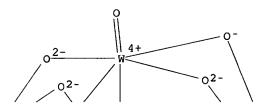
Absolute stereochemistry.

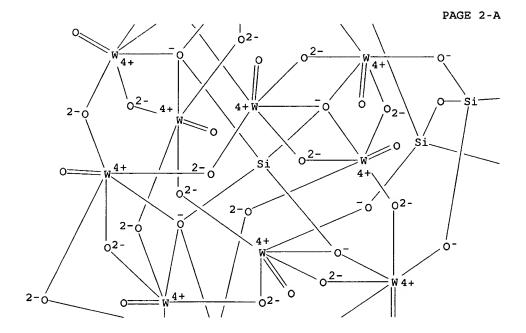
RN 215594-72-2 HCAPLUS

CN Methanaminium, N,N,N-trimethyl-, [μ4-[1,3-bis(3-chloropropyl)1,1,3,3-disiloxanetetrolato(4-)-κO1:κO1':κO3:.ka
ppa.O3']] [μ11-[orthosilicato(4-)-κO:κO:κO:.ka
ppa.O':κO':κO':κO'':κO'':κO'':.kappa
.O''':κO''']]eicosa-μ-oxoundecaoxoundecatungstate(4-)
(4:1) (9CI) (CA INDEX NAME)

CM 1

CRN 215594-71-1 CMF C6 H12 Cl2 O40 Si3 W11 CCI CCS

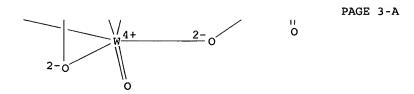




PAGE 2-B

$$---- (CH2)3 - C1$$

$$\sim$$
 (CH₂)₃-Cl



CM 2

CRN 51-92-3 CMF C4 H12 N

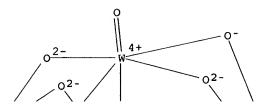
RN 215594-74-4 HCAPLUS

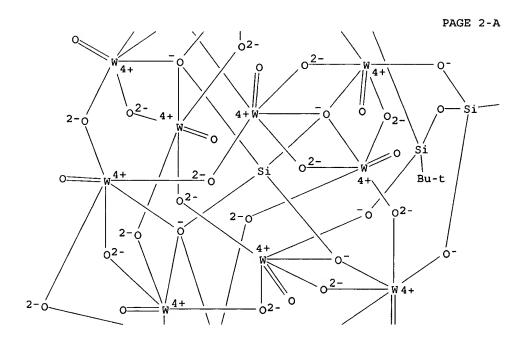
CN Methanaminium, N,N,N-trimethyl-, [μ 4-[1,3-bis(1,1-dimethylethyl)-1,1,3,3-disiloxanetetrolato(4-)- κ 01: κ 01': κ 03: κ 03']] [μ 11-[orthosilicato(4-)- κ 0: κ 0': κ 0': κ 0': κ 0'': κ 0'': κ 0'': κ 0'': κ 0'': κ 0'': κ 0''': κ 0'''|]]eicosa- μ -oxoundecaoxoundecatungstate(4-) (4:1) (9CI) (CA INDEX NAME)

CM 1

CRN 215594-73-3 CMF C8 H18 O40 Si3 W11 CCI CCS

PAGE 1-A





PAGE 2-B

2-0 W 4+ 2-0 O O

CM 2

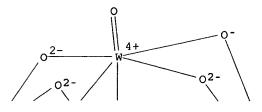
CRN 51-92-3 CMF C4 H12 N

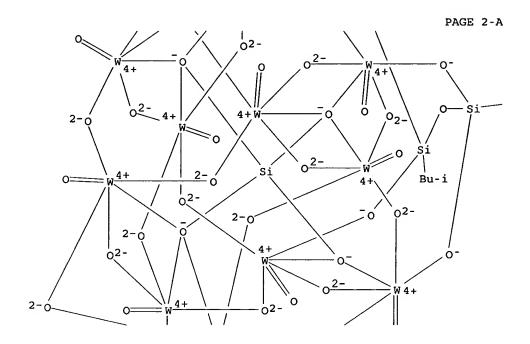
CCI CCS

RN 215594-76-6 HCAPLUS
CN Methanaminium, N,N,N-trimethyl-, [μ4-[1,3-bis(2-methylpropyl)1,1,3,3-disiloxanetetrolato(4-)-κO1:κO1':κO3:.ka
ppa.O3']] [μ11-[orthosilicato(4-)-κO:κO:κO:.ka
ppa.O':κO':κO':κO'':κO'':κO'':.kappa
.O''':κO''']]eicosa-μ-oxoundecaoxoundecatungstate(4-)
(4:1) (9CI) (CA INDEX NAME)

CM 1

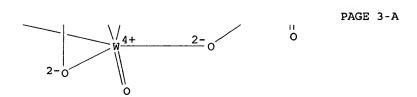
CRN 215594-75-5
CMF C8 H18 O40 Si3 W11





PAGE 2-B

— Ви-і

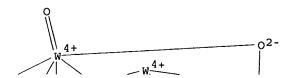


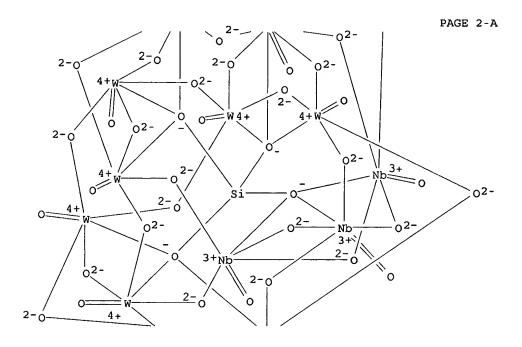
CM 2

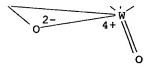
CRN 51-92-3 CMF C4 H12 N

RN

215594-80-2 HCAPLUS Niobate(7-), [µ12-[orthosilicato(4-)-CN

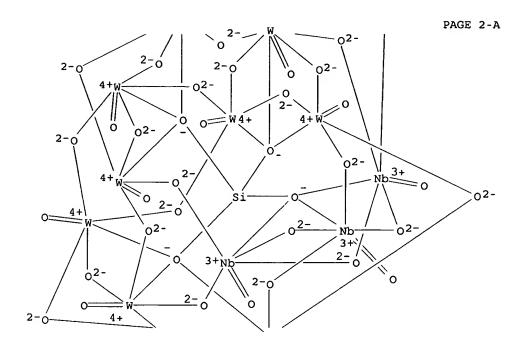
Les Henderson Page 291 571-272-2538 

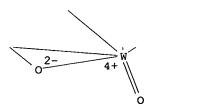




●7 Cs+





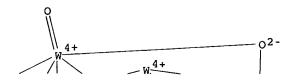


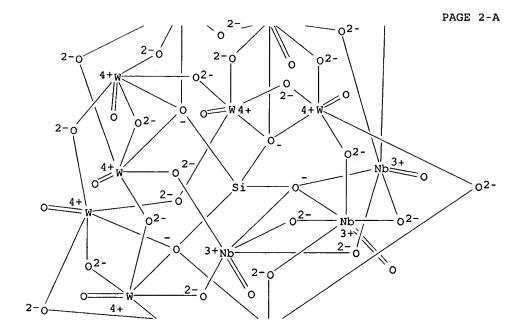
●7 H+

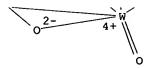
CM 2

CRN 113-00-8 CMF C H5 N3

 $\begin{array}{c} \text{NH} \\ || \\ \text{H}_2 \text{N--} \text{C---} \text{NH}_2 \end{array}$

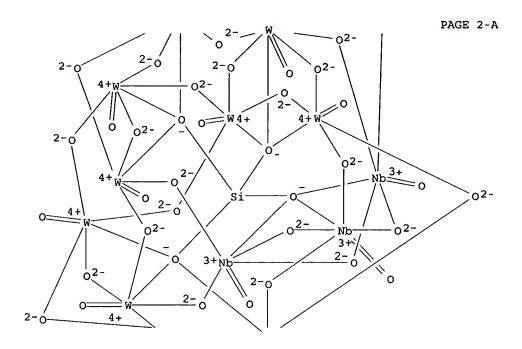


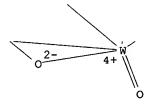




●7 Rb+







●7 H+

CM 2

CRN 110-86-1 CMF C5 H5 N



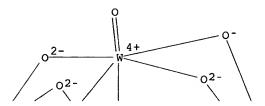
CM 1

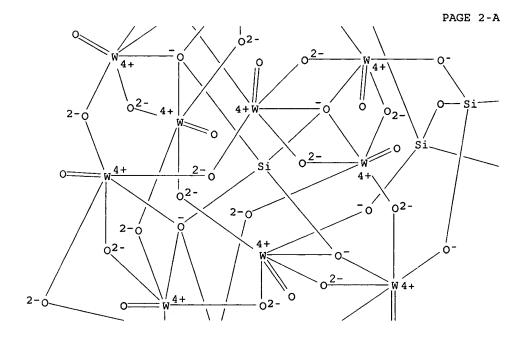
CRN 215594-85-7

CMF C32 H66 O40 Si3 W11

CCI CCS

PAGE 1-A

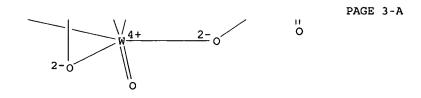




PAGE 2-B

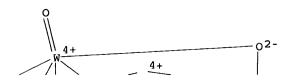
$$----$$
 (CH₂)₁₅-Me

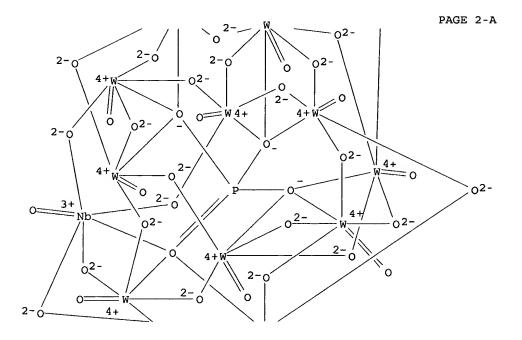
 \sim (CH₂)₁₅-Me

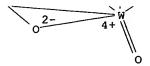


CM 2

.CRN 51-92-3 CMF C4 H12 N

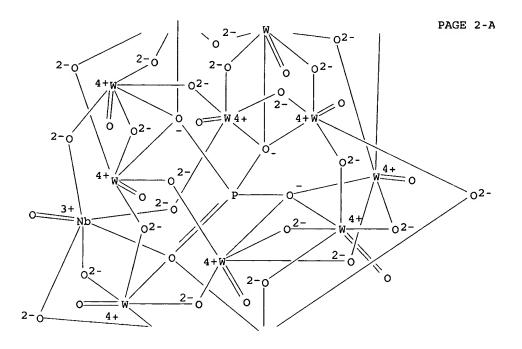


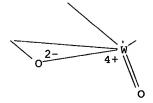




●4 K+

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





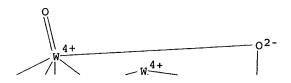
PAGE 3-A

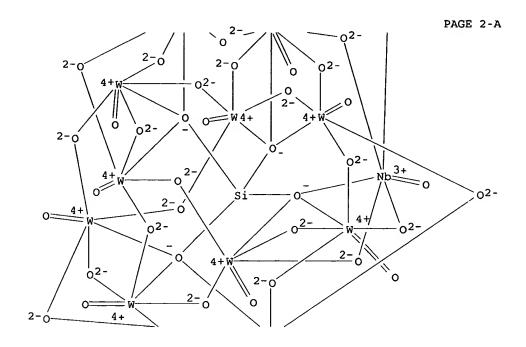
●4 H+

CM 2

CRN 75-50-3 CMF C3 H9 N

RN 215594-93-7 HCAPLUS
CN Niobate(5-), (eicosa-μ-oxoundecaoxoundecatungstate) [μ12[orthosilicato(4-)-κΟ:κΟ:κΟ':κΟ':
κΟ':κΟ'':κΟ'':κΟ''':κΟ'''
:κΟ'''] tetra-μ-oxooxo-, pentapotassium (9CI) (CA INDEX NAME)



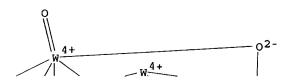


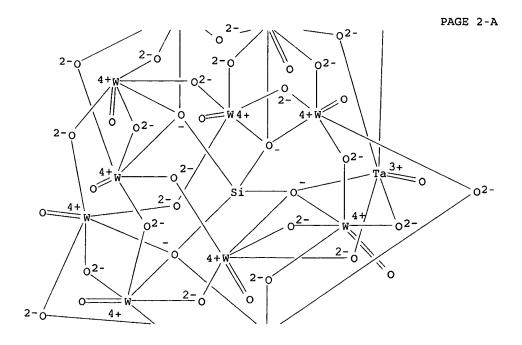


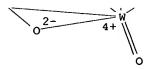
●5 K+

RN 215594-95-9 HCAPLUS CN Tantalate(5-), (eicosa- μ -oxoundecaoxoundecatungstate) [μ 12-[orthosilicato(4-)- κ 0: κ 0: κ 0': κ 0'

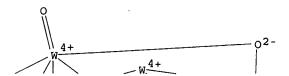
PAGE 1-A

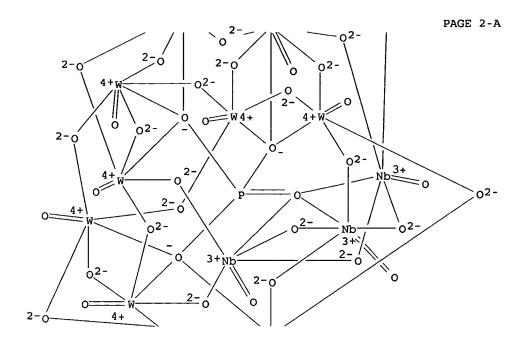


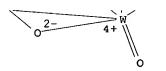




●5 K+

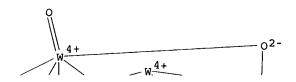


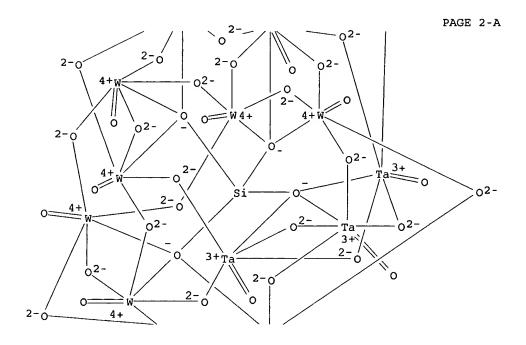


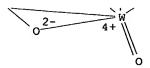


●6 K+

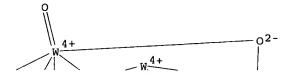
PAGE 1-A

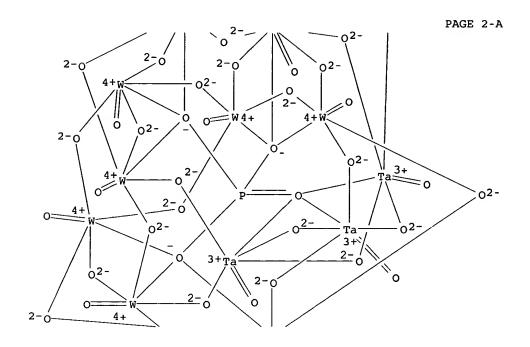


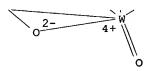




●7 K+







●6 K+

RN 215595-09-8 HCAPLUS
CN Methanaminium, N,N,N-trimethyl-, hydrogen cobaltate[µ4-[1,3-diethyl-1,1,3,3-disiloxanetetrolato(4-)κO1:κO1':κO3:κO3']]eicosa-µ-oxo-µ3oxotri-µ4-oxoundecaoxoundecatungstate(6-) (4:2:1) (9CI) (CA INDEX NAME)

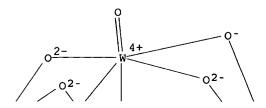
CM 1

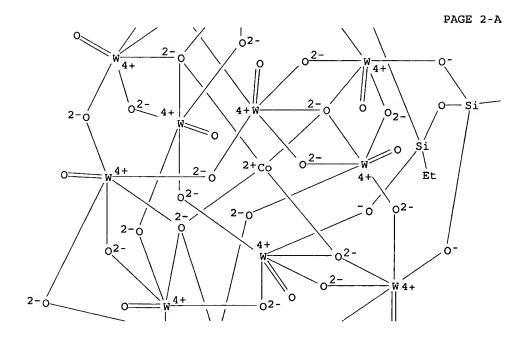
CRN 215595-08-7

CMF C4 H10 Co O40 Si2 W11

CCI CCS

PAGE 1-A





PAGE 2-B

PAGE 3-A 2-0

CM 2

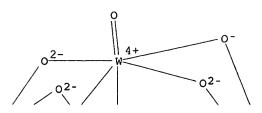
CRN 51-92-3 CMF C4 H12 N

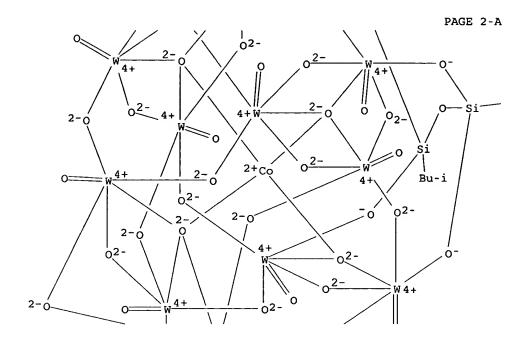
RN 215595-11-2 HCAPLUS Methanaminium, N,N,N-trimethyl-, dihydrogen [μ4-[1,3-bis(2-CN methylpropyl)-1,1,3,3-disiloxanetetrolato(4-)- $\kappa01:\kappa01':\kappa03:\kappa03']$]cobaltateeicosa- μ -

 $oxo-\mu3-oxotri-\mu4-oxoundecaoxoundecatungstate(6-)$ (4:2:1) (9CI) (CA INDEX NAME)

CM 1

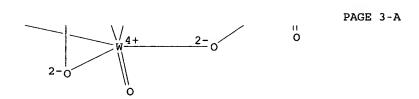
CRN 215595-10-1 CMF C8 H18 Co O40 Si2 W11 CCI CCS





PAGE 2-B

— Ви-і



CM 2

CRN 51-92-3 CMF C4 H12 N

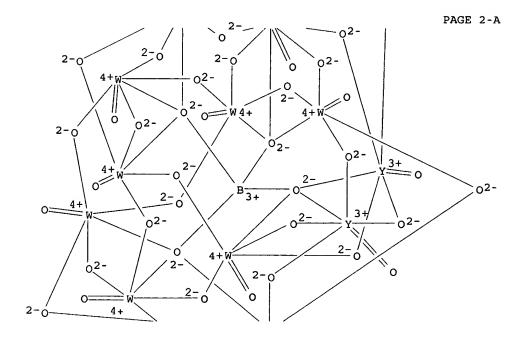
RN

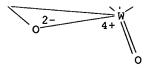
215595-19-0 HCAPLUS Yttrate(7-), (heptadeca- μ -oxodecaoxodecatungstate)hepta- μ -oxodioxo[μ 12-[tetrahydroxyborato(5-)-CN

κΟ:κΟ:κΟ':κΟ':κΟ':κΟ' ':κΟ'':κΟ'':κΟ''':κΟ''':κΟ''']]di-, heptapotassium (9CI) (CA INDEX NAME)

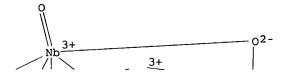
PAGE 1-A

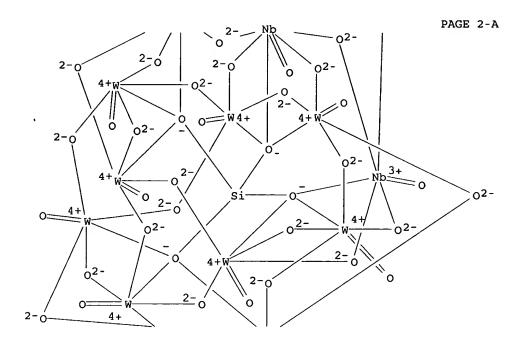


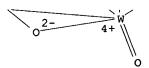




●7 K+







K+

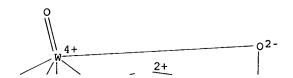
215595-22-5 HCAPLUS RNCNL-Lysine, (heptadeca- μ -oxodecaoxodecatungstate)hepta- μ oxodioxo[μ12-[phosphato(3-)-κ0:κ0:κ0:κ0':κ0':κ0':κ0'':κ0'':κ0'' :κΟ''':κΟ''']]dititanate(7-) (7:1) (9CI) (CA INDEX NAME)

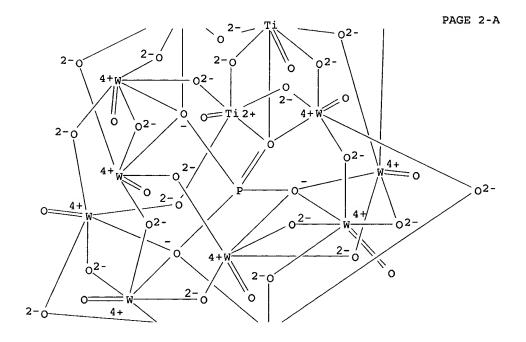
CRN 215601-31-3 CMF H . 1/7 O40 P Ti2 W10 CCI CCS

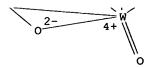
1

CM

PAGE 1-A







●7 H+

CM 2

CRN 56-87-1 CMF C6 H14 N2 O2

Absolute stereochemistry.

RN 215595-24-7 HCAPLUS

CN Tungstate(6-), tetracosa- μ -oxotetra- μ 4oxododecaoxozincatedodeca-, hexahydrogen (9CI) (CA INDEX NAME)

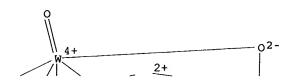
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

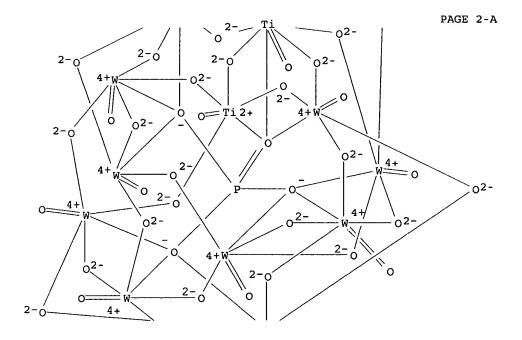
RN 215601-32-4 HCAPLUS

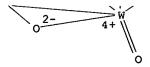
CN L-Histidine, (heptadeca-µ-oxodecaoxodecatungstate)hepta-µoxodioxo[μ 12-[phosphato(3-)- κ 0: κ 0: κ 0: κ 0 ':κ0':κ0':κ0'':κ0'':κ0'':κ0''' $\kappa O''' \kappa O'''$]dititanate(7-) (7:1) (9CI) (CA INDEX NAME)

1 CM

CRN 215601-31-3 CMF H . 1/7 O40 P Ti2 W10 CCI CCS







●7 H1

CM 2

CRN 71-00-1 CMF C6 H9 N3 O2

Absolute stereochemistry. Rotation (-).

IC ICM A01N055-02

ICS A01N059-14; A01N059-16; A01N059-20

INCL 514492000

CC 1-5 (Pharmacology)

Section cross-reference(s): 29, 63, 78

ST respiratory viral infection antiviral aerosol

polyoxometalate

IT Respiratory tract

(infection, viral; method and aerosol spray containing a polyoxometalate for treating and preventing respiratory viral infections)

IT Human immunodeficiency virus 1

Orthomyxovirus

Paramyxovirus

(inhibition of; method and aerosol spray containing a polyoxometalate for treating and preventing respiratory viral infections)

IT Antiviral agents

Influenza A virus

Influenza B virus

Respiratory syncytial virus

(method and aerosol spray containing a polyoxometalate for treating and preventing respiratory viral infections)

IT Heteropoly acids

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(method and aerosol spray containing a polyoxometalate for treating and preventing respiratory viral infections)

IT Erythrocyte

(polyoxometalate inhibition of influenza A virus-caused hemagglutination and hemolysis of, of chick; method and aerosol spray containing a polyoxometalate for treating and preventing respiratory viral infections)

IT Hemagglutination

Hemolysis

(polyoxometalate inhibition of influenza A virus-caused, of chick erythrocytes; method and aerosol spray

containing a polyoxometalate for treating and preventing



```
respiratory viral infections)
IT
     Drug delivery systems
        (sprays; method and aerosol spray containing a
        polyoxometalate for treating and preventing respiratory
        viral infections)
IT
     Infection
        (viral, respiratory; method and aerosol spray containing a
        polyoxometalate for treating and preventing respiratory
        viral infections)
IT
     11078-54-9 12027-38-2D, solid solution with
     ammonium analog 12045-18-0 12059-48-2 12141-67-2
     12142-54-0
                12200-88-3 12297-12-0 12297-12-0D
     , solid solns. with ammonium analog and protonated amino acid
     analog 12329-09-8 12329-10-1 12390-22-6 12411-74-4
     12436-83-8 37300-95-1 39282-41-2
                                            59054-50-1 59111-46-5
     63950-53-8 63995-70-0
                              64684-58-8
                                            70316-17-5
                                                          75656-59-6
     77981-80-7D, solid solution with tetrahydrogen analog
     79104-95-3 81552-97-8 82679-05-8
     83721-03-3 83721-04-4 84303-03-7
     84303-05-9
                  84750-84-5 87261-30-1
                                           89899-81-0,
    Ammonium antimony sodium tungsten oxide ((NH4)17Sb9Na2W21086)
     92762-45-3 92767-45-8 93425-27-5 100513-52-8
     101144-77-8 101346-99-0 101347-00-6 101347-04-0
     101347-05-1 101347-09-5 101347-11-9
                                                101347-12-0
    101347-13-1 102073-48-3 108834-36-2 108987-13-110294-54-7 110313-16-1 110717-64-1 110717-65-110717-67-4 110717-70-9 111933-31-4 112763-08-3
                                                108987-13-9
                                                110717-65-2
     112763-08-3D, solid solution with tetrahydrogen analog
    116434-67-4 119390-04-4 119720-71-7 119923-89-6 123639-37-2 129238-68-2 129238-69-3D, solid solns. with sodium
     and tetramethylammonium analogs 129238-70-6D, solid solns. with
     sodium and tetramethylammonium analogs 129572-46-9
    129572-47-0 129592-85-4 131359-48-3
     131541-68-9 131541-69-0 131541-70-3
     132460-56-1 132460-57-2 132460-58-3
     134107-05-4 138026-47-8 139631-90-6 139631-92-8
     139631-93-9 139631-95-1 139631-96-2
    139631-98-4 139632-00-1 141483-63-8
    141532-40-3 141532-61-8 143823-91-0 143823-92-1
    144547-23-9 146026-67-7 148362-93-0 149275-00-3
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153541-07-2
                                              152514-03-9
158702-61-5
     152444-38-7
                  152444-39-8
    153481-12-0 153481-15-3
    160097-69-8D, solid solns. with sodium and potassium analogs
    160097-70-1D, solid solns. with potassium and sodim analogs
    160220-13-3
                 160241-96-3 162958-07-8D, solid solns. with
    potassium and tetramethylammonium analogs 162958-09-0
    162958-11-4 162958-12-5 162958-14-7 162958-16-9 162958-18-1 162958-20-5
                                             162958-21-6
    tetramethylammonium analogs 163128-97-0 163128-98-1
    163151-26-6 163151-27-7 167397-05-9 170126-82-6
    189823-33-4
    189823-37-8 189823-38-9 194150-76-0 215545-74-7
    215545-75-8, Potassium titanium tungsten oxide (K7Ti2W10O40)
    215545-77-0 215545-78-1 215545-79-2 215545-80-5
    215545-81-6 215545-82-7 215545-83-8 215545-84-9
215545-85-0 215545-86-1 215591-58-5 215594-35-7
215594-50-6 215594-63-1 215594-64-2 215594-65-3D,
    solid solution with histidine or lysine analog 215594-66-4D
     , solid solution with sodium analog
                                           215594-68-6
                                                        215594-70-0
    215594-72-2 215594-74-4 215594-76-6
    215594-77-7 215594-78-8 215594-79-9 215594-80-2
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    215594-86-8 215594-87-9 215594-88-0 215594-89-1
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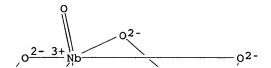
CN

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     215595-12-3 215595-13-4 215595-14-5
                                               215595-15-6
     215595-16-7 215595-17-8D, Niobium sodium tungsten oxide
     (Nb4Na6W2O19), solid solution with potassium analog 215595-18-9
     215595-19-0 215595-21-4 215595-22-5D,
     solid solution with sodium analog 215595-23-6D, Niobium potassium
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     215595-24-7 215601-32-4 215601-56-2
     215601-59-5
     RL: BAC (Biological activity or effector, except adverse); BSU
     (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (method and aerosol spray containing a polyoxometalate
        for treating and preventing respiratory viral infections)
REFERENCE COUNT:
                         100
                               THERE ARE 100 CITED REFERENCES AVAILABLE
                               FOR THIS RECORD. ALL CITATIONS AVAILABLE
                               IN THE RE FORMAT
L114 ANSWER 40 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
                         1997:383186 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         127:109043
TITLE:
                         Syntheses and characterization of the
                         heptasodium salt of the Keggin-type
                         triniobium-substituted polyoxoanion
                         SiW9Nb307-40 and the all-sodium salt of the
                         polyoxoanion-supported organometallic complex
                         [(n5-C5Me5)Rh·SiW9Nb3O40]5-
AUTHOR(S):
                         Nomiya, Kenji; Nozaki, Chika; Kano, Atsuyuki;
                         Taguchi, Takayuki; Ohsawa, Katsunori
                         Department of Materials Science, Kanagawa
CORPORATE SOURCE:
                         University, Hiratsuka, Kanagawa, 259-12, Japan
                         Journal of Organometallic Chemistry (1997),
SOURCE:
                         533(1-2), 153-159
                         CODEN: JORCAI; ISSN: 0022-328X
PUBLISHER:
                         Elsevier
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     The prepns. of A-\beta-Na7SiW9Nb3040 and Na5[(\eta5-
     C5Me5)Rh·SiW9Nb3O40] are described. The water-soluble form of
     the Keggin-type trisubstituted heteropolyanion,
    A-\beta-SiW9Nb307-40, of interest as a polyoxoanion support for
    organometallic complexes, was isolated in pure form in
     61% yield as its heptasodium salt with 16 waters of hydration.
     The polyoxoanion-supported organometallic complex as its
     all-sodium salt, [(\eta 5\text{-C5Me5})Rh·SiW9Nb3O40]5- was
    obtained in 21% yield as an anal. pure, homogeneous
    yellow-brown solid by the reaction of [(η5-C5Me5)Rh(CH3CN)3]2+
    with the trisubstituted heteropolyanion SiW9Nb307-40 in
    CH3CN-DMSO. These all-sodium salts have required their own novel
    prepns. and purifications, and have never been derived
     directly from the known all-Bu4N+ salts of the corresponding
    Keggin heteropolyanions. Compositional characterization
    was accomplished by complete elemental analyses, TG/DTA, and
    FT-IR. Structural characterization in solution was achieved by
    combination of 1H, 13C and 183W NMR spectroscopies.
    192314-19-5P 192314-22-0P
    RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of)
    192314-19-5 HCAPLUS
    Niobate (7-), [\mu 12-[orthosilicato (4-)-
    κ0:κ0:κ0:κ0':κ0':κ0':κ0'
     ':κ0'':κ0'':κ0''':κ0''':κ0''']]nona-
     μ-oxotrioxo (pentadeca-μ-oxononaoxononatungstate) tri-,
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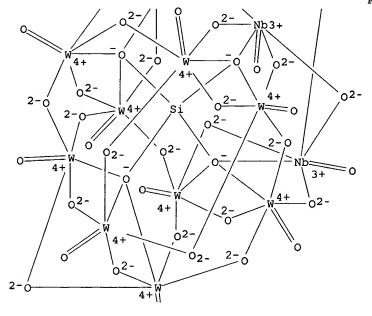
Les Henderson Page 321 571-272-2538

heptasodium, hexadecahydrate (9CI) (CA INDEX NAME)

PAGE 1-A



PAGE 2-A



0

●7 Na+

●16 H₂O

RN 192314-22-0 HCAPLUS
CN Niobate(5-), [μ12-[orthosilicato(4-)κΟ:κΟ:κΟ:κΟ':κΟ':κΟ':κΟ'
':κΟ':κΟ':κΟ':κΟ':κΟ'']]heptaμ-oxotri-μ3-oxotrioxo[[(1,2,3,4,5-η)-1,2,3,4,5pentamethyl-2,4-cyclopentadien-1-yl]rhodate](tetradeca-μoxononaoxononatungstate)tri-, pentasodium, compd. with
sulfinylbis[methane] (1:3), dihydrate (9CI) (CA INDEX NAME)

CM 1

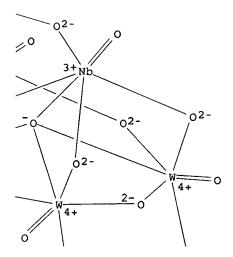
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CCI CCS

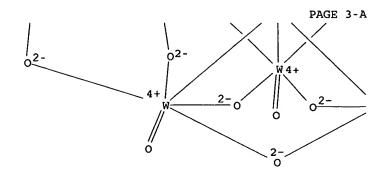
PAGE 1-A

Ме

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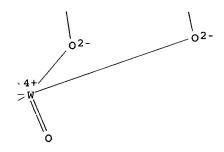
PAGE 2-B





●5 Na+

PAGE 3-B



CM 2

CRN 67-68-5 CMF C2 H6 O S

CC 29-13 (Organometallic and Organometalloidal Compounds) Section cross-reference(s): 78

ST triniobium nonatungsten polyoxoanion Keggin type prepn; rhodium triniobium nonatungsten polyoxoanion Keggin type; tungsten rhodium sodium niobium polyoxometallate Keggin

ΙT 192314-19-5P 192314-22-0P

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

REFERENCE COUNT:

19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L114 ANSWER 41 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1997:375889 HCAPLUS

DOCUMENT NUMBER: 127:8415

TITLE: New reactive carbons for HD decomposition AUTHOR(S): Walker, John E.; Rivin, Donald; Kendrick, Cyrus E.; Hill, Craig L.

CORPORATE SOURCE:

SOURCE:

Development Engineering Center, U. S. Army Natick Research, Natick, MA, 01760-5020, USA Proceedings of the ERDEC Scientific Conference on Chemical and Biological Defense Research, Aberdeen Proving Ground, Md., Nov. 15-18, 1994 (1996), Meeting Date 1994, 821-828.

Editor(s): Berg, Dorothy A. National Technical Information Service: Springfield,

Va.

CODEN: 64NAAX Conference

DOCUMENT TYPE: LANGUAGE:

English

In this paper we report on the reactivity of HD simulants by catalytic carbon systems formulated by the incorporation of the heteropolyoxometalate, H5PV2Mo10040 (POX) into microporous carbon. The activated carbon systems are; Ambersorb (Rohm & Haas), Maxsorb (Kansai) and Calgon PCB-G (Calgon Corp.). ESR studies indicate that the POX is strongly adsorbed to the surface of the carbons. Uing chloroethyl ethylsulfide (CEES) as an HD simulant, we found that the POX-carbons oxidized CEES in both acetonitrile and toluene solvent systems. GC/MS analyses of the reaction products showed only the presence of the sulfoxide of CEES with no evidence of the sulfone. Static vapor testing of the POX-carbons with CEES showed that the reactivity varies as a function of the water vapor introduced with the CEES vapor. Microflow calorimetry of the POX-carbons using trichloroethylene (TCE) as the adsorptive showed no appreciable decrease in the heats of adsorption and desorption when compared to the carbon controls. When the HD simulant tetrahydrothiophene (THT) was used as the adsorptive, there was evidence of a chemisorption reaction occurring along with the phys. adsorption process. incorporation of polyoxometalates into microporous carbon supports should lead to a new generation of sorptive/reactive chemical protective materials.

IT 12293-21-9

CN

RL: CAT (Catalyst use); USES (Uses)

(new reactive carbons for HD decomposition)

RN 12293-21-9 HCAPLUS

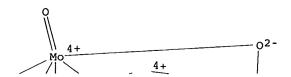
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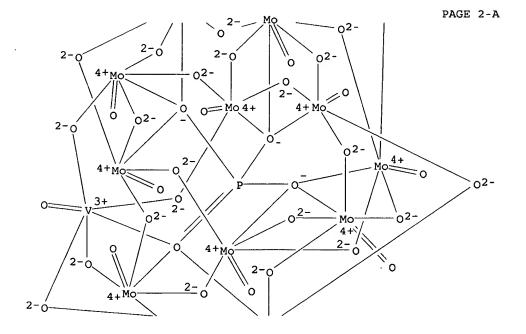
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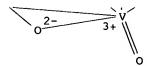
 $:\kappa O''':\kappa O''']$ di-, pentahydrogen (9CI) (CA INDEX

NAME)

PAGE 1-A







●5 H+

CC 59-4 (Air Pollution and Industrial Hygiene)

Section cross-reference(s): 50

12293-21-9 TT

RL: CAT (Catalyst use); USES (Uses)

(new reactive carbons for HD decomposition)

L114 ANSWER 42 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

1996:737410 HCAPLUS ACCESSION NUMBER:

126:93902 DOCUMENT NUMBER:

The interaction of nitrogen oxides with TITLE:

metal-oxygen cluster compounds (heteropoly

oxometalates)

AUTHOR(S): Belanger, R.; Moffat, J. B.

CORPORATE SOURCE: Department of Chemistry and the

Guelph-Waterloo Centre for Graduate Work in Chemistry, University of Waterloo, Waterloo,

Ontario, Can.

SOURCE: Journal of Molecular Catalysis A: Chemical

(1996), 114(1-3), 319-329 CODEN: JMCCF2; ISSN: 1381-1169

PUBLISHER: Elsevier DOCUMENT TYPE: Journal LANGUAGE: English

NO2 is sorbed by the solid heteropoly acids, the 1st sorbates producing HNO3 while those subsequently taken up remain strongly bound on the acid, ≤3 NO2/heteropoly anion. The sorption of NO2 depends on the elemental composition of the anions of the solid acids and hence on the acid strengths of these materials. The sorbed NO2 assocs. with both the surface and bulk protons to form HNO2+ thus demonstrating that NO2 is capable of penetrating into the crystallog. structure of the solid acid. Prior sorption of NO2 facilitates the takeup of NO which forms N2O3; 12-tungstophosphoric acid (HPW) supported on silica produces quant. different but qual. similar results. The microporous Al salt of HPW produces markedly different results when exposed to NO2 with the predominant product being N evidently resulting from the ammonium cation functioning as a source of the reductant ammonia.

1343-93-7, 12-Tungstophosphoric acid 12027-38-2, TT

12-Tungstosilicic acid

RL: NUU (Other use, unclassified); USES (Uses)

(interaction of nitrogen oxides with metal-oxygen cluster compds.)

RN 1343-93-7 HCAPLUS

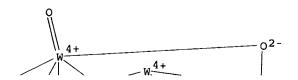
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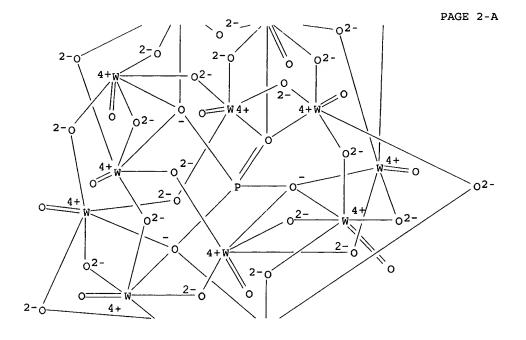
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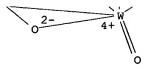
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a-, trihydrogen (9CI) (CA INDEX NAME)

PAGE 1-A

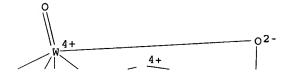


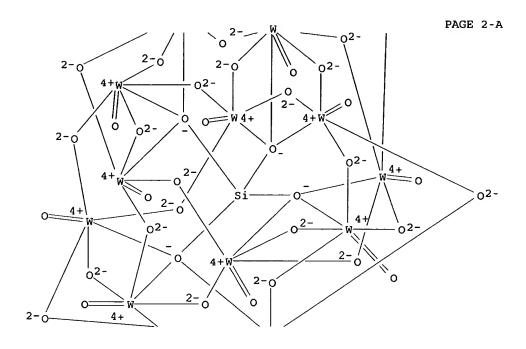


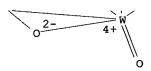


●3 H+

PAGE 1-A







●4 H+

CC 59-4 (Air Pollution and Industrial Hygiene)

Section cross-reference(s): 67

IT 1343-93-7, 12-Tungstophosphoric acid 12027-38-2,

12-Tungstosilicic acid

RL: NUU (Other use, unclassified); USES (Uses)

(interaction of nitrogen oxides with metal-oxygen cluster

compds.)

REFERENCE COUNT:

61 THERE ARE 61 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE

IN THE RE FORMAT

L114 ANSWER 43 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1996:410483 HCAPLUS

DOCUMENT NUMBER:

125:49278

TITLE:

Method, compositions, and apparatus

for treating and preventing respiratory viral

infections with polyoxometalate

aerosol

Patent

INVENTOR (S):

Schinazi, Raymond F.; Hill, Craig L.

PATENT ASSIGNEE(S): USA

SOURCE:

PCT Int. Appl., 71 pp.

CODEN: PIXXD2

DOCUMENT TYPE: LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 2 PATENT INFORMATION:

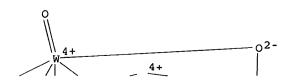
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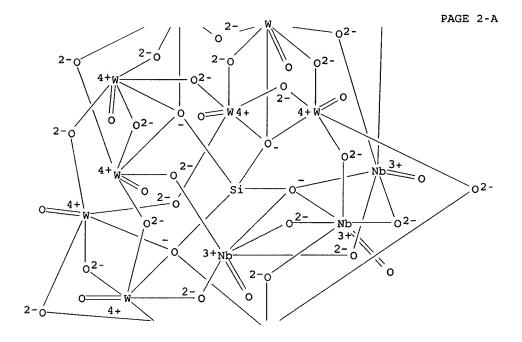
CRN 162958-08-9 CMF H . 1/7 Nb3 O40 Si W9 CCI CCS

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
	O 9609764	Al	19960404	WO 1995-US11961	1995 0926	
	ES, FI, GB, LT, LU, LV, RU, SD, SE,	GE, HU, MD, MG, SG, SI,	IS, JP, K MK, MN, M SK, TJ, T	CA, CH, CN, CZ, DE, DK, CE, KG, KP, KR, KZ, LK, W, MX, NO, NZ, PL, PT, M	EE, LR, RO,	
**	IE, IT, LU, GN, ML, MR,	MC, NL, NE, SN,	PT, SE, B	F, BJ, CF, CG, CI, CM,		
	IS 5824706	A	19981020		1995 0303	
A	.U 9536366	A1	19960419	AU 1995-36366	1995 0926	
PRIORI	TY APPLN. INFO.:			US 1994-312561	A 1994 0926	
				US 1995-399700	A 1995 0303	
				WO 1995-US11961	W 1995 0926	
AB Respiratory viral infections may be effectively prevented or treated by administering an aerosol spray comprising a polyoxometalate to the lungs. More than 200 polyoxometalates are claimed. The virus may be e.g. influenza A, influenza B, HIV-1, or respiratory syncytial virus.						
IT 162958-09-0 RL: ADV (Adverse effect, including toxicity); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (polyoxometalate aerosols for treating and preventing						
CN N O o h						

Les Henderson Page 332 571-272-2538

PAGE 1-A





₽7 H+

CM 2 CRN 75-50-3 CMF C3 H9 N

CH₃ H3C-N-CH3

IC ICM A01N055-02 ICS A01N059-14; A01N059-16; A01N059-20; A61K031-28; A61K031-295; A61K031-30; A61K031-315; A61K033-22; A61K033-24; A61K033-26; A61K033-32; A61K033-34

1-5 (Pharmacology) CC

Section cross-reference(s): 63

ST polyoxometalate aerosol respiratory virus infection; antiviral respiratory virus polyoxometalate aerosol

Virucides and Virustats ΙT

(polyoxometalate aerosols for treating and preventing respiratory viral infections)

IT Heteropoly acids

> RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (polyoxometalate aerosols for treating and preventing respiratory viral infections)

TT Respiratory tract

(disease, infection, polyoxometalate aerosols for

treating and preventing respiratory viral infections)

IT Virus, animal

(human immunodeficiency 1, polyoxometalate aerosols for treating and preventing respiratory viral infections)

Virus, animal TΤ

(influenza A, polyoxometalate aerosols for treating and preventing respiratory viral infections)

IT Virus, animal

(influenza B, polyoxometalate aerosols for treating and preventing respiratory viral infections)

TT Virus, animal

(measles, polyoxometalate aerosols for treating and preventing respiratory viral infections)

TΤ Virus, animal

(mumps, polyoxometalate aerosols for treating and preventing respiratory viral infections)

TΨ Virus, animal

(parainfluenza 2, polyoxometalate aerosols for treating and preventing respiratory viral infections)

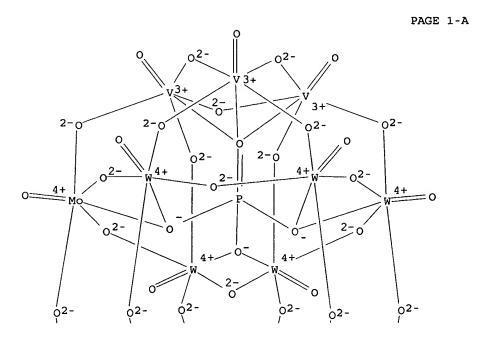
TΤ Virus, animal

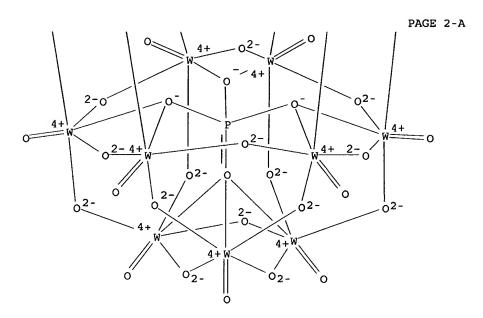
(parainfluenza 3, polyoxometalate aerosols for treating and preventing respiratory viral infections)

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IT
     Virus, animal
         (respiratory syncytial, polyoxometalate aerosols for
         treating and preventing respiratory viral infections)
     Pharmaceutical dosage forms
ΙT
         (sprays, polyoxometalate aerosols for treating and
        preventing respiratory viral infections)
     84750-84-5 162958-09-0
     RL: ADV (Adverse effect, including toxicity); BAC (Biological
     activity or effector, except adverse); BSU (Biological study,
     unclassified); THU (Therapeutic use); BIOL (Biological study);
     USES (Uses)
         (polyoxometalate aerosols for treating and preventing
         respiratory viral infections)
L114 ANSWER 44 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 1995:774639 HCAPLUS
DOCUMENT NUMBER:
                           123:160817
TITLE:
                          Polyoxometallates in the treatment
                           of flavivirus infections
INVENTOR(S):
                           Weigold, Helmut
                           Commonwealth Scientific and Industrial
PATENT ASSIGNEE(S):
                           Research Organization, Australia
SOURCE:
                           PCT Int. Appl., 36 pp.
                           CODEN: PIXXD2
DOCUMENT TYPE:
                           Patent
LANGUAGE:
                           English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                      KIND DATE
                                             APPLICATION NO.
                                                                       DATE
                                  -----
                                               -----
                          ----
                      A1 19950427 WO 1994-AU641
     WO 9511033
                                                                        1994
         W: AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, JP, KE, KG, KP, KR, KZ, LK, LR, LT, LU, LV, MD, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SI, SK, TJ, TT, UA, US, UZ
         RW: KE, MW, SD, SZ, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE,
              IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN,
              ML, MR, NE, SN, TD, TG
                           A1
                                  19950508
                                             AU 1994-79853
     AU 9479853
                                                                        1994
                                                                        1021
PRIORITY APPLN. INFO.:
                                               AU 1993-1950
                                                                        1993
                                                                        1022
                                               WO 1994-AU641
                                                                        1994
                                                                        1021
     Pharmaceutical compns. containing polyoxometallates
AB
     and pharmaceutically acceptable derivs. thereof are disclosed.
     Also disclosed is the use of such compds. or compns. in
     therapy for the treatment or prophylaxis of infections by viruses
     which are confirmed or probable members of the family
     Flaviviridae, including Hepatitis C. In an assay using type 2
     dengue virus-infected Vero cells, K6[P2V2W16O62]·40H2O had
     an inhibitory concentrate of 5-10 μM.
     110390-83-5P 133348-29-5P 133348-30-8P
IT
     167308-24-9P
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RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation);

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USES (Uses)
         (polyoxometallates for treatment of flavivirus
         infections)
     110390-83-5 HCAPLUS
RN
     Vanadate (7-), [heptacosa-\mu-oxopentadecaoxo[\mu9-[phosphato(3-)-
CN
     κ0:κ0:κ0:κ0':κ0':κ0'':κ0
     '':κ0''':κ0''']]pentadecatungstate]octa-μ-
     oxooxo(\mu-oxodioxodimolybdate)[\mu9-[phosphato(3-)-
     κ0:κ0:κ0:κ0':κ0':κ0'':κ0
     '':κO''':κO''']]-, heptapotassium (9CI) (CA INDEX
     NAME)
    STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN
     133348-29-5 HCAPLUS
     Vanadate (8-), [heptacosa-\mu-oxopentadecaoxo[\mu9-phosphato(3-)-0:0:0':0':0'':0'':0''']-pentadecatungstate]nona-\mu-
CN
     oxodioxo(oxomolybdate)[µ9-[phosphato(3-)-
     0:0:0:0':0':0'':0'':0''':0''']]di-, octapotassium (9CI) (CA INDEX
     NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN
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CN
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     '':κ0''':κ0''']]hexadecatungstate]hepta-μ-
     oxodioxo [μ9-[phosphato(3-)-κ0:κ0:κ0:κ0
     :κ0':κ0'':κ0''':κ0''']]di-,
octapotassium (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     167308-24-9 HCAPLUS
RN
CN
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     oxotetradecaoxo[µ9-[phosphato(3-)-0:0:0:0':0':0':0'':0'':0'''
     ]]tetradecatungstate]tri-, nonapotassium (9CI) (CA INDEX NAME)
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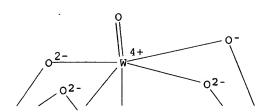


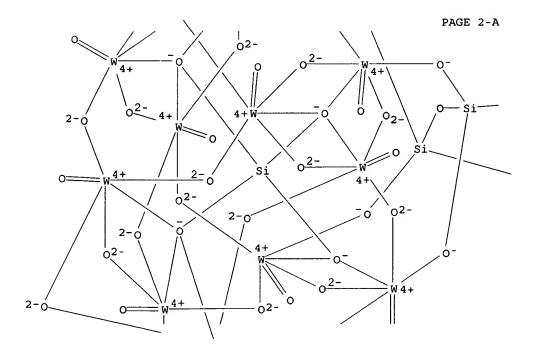


●9 K+

IT 139631-93-9P
 RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL
 (Biological study); PREP (Preparation); USES (Uses)
 (polyoxometallates for treatment of flavivirus
 infections)
RN 139631-93-9 HCAPLUS
CN Tungstate(4-), [μ11-[orthosilicato(4-) κΟ:κΟ:κΟ:κΟ':κΟ':κΟ':κΟ'
 ':κΟ':κΟ':κΟ':κΟ':']]eicosa-μ oxoundecaoxo[μ4-[[4,4'-[1,1,3,3-tetra(hydroxy-κΟ)-1,3-disiloxanediyl]bis[butanenitrilato]](4-)]]undeca-, tetracesium
 (9CI) (CA INDEX NAME)

PAGE 1-A





* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

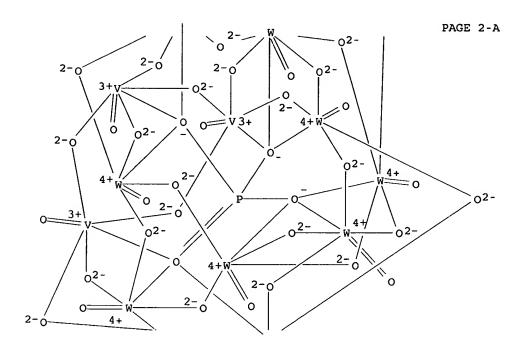
●4 Cs+

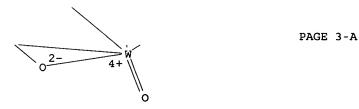
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ICM A61K033-24
IC
CC
     1-5 (Pharmacology)
    Section cross-reference(s): 63, 78
    polyoxometallate flavivirus infection treatment
ST
     Pharmaceutical dosage forms
IT
     Virucides and Virustats
        (polyoxometallates for treatment of flavivirus
        infections)
TT
     Virus, animal
        (dengue 2, polyoxometallates for treatment of
        flavivirus infections)
IT
     Virus, animal
        (flavi-, polyoxometallates for treatment of
        flavivirus infections)
IT
     Heteropoly acids
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (tungstates, polyoxometallates for treatment of
        flavivirus infections)
                                               133348-21-7P
TΤ
    110313-16-1P 110390-83-5P
                                111933-31-4P
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                                139381-06-9P
     167308-24-9P 167308-27-2P
                                  167308-29-4P
                                                167308-30-7P
                  167308-32-9P
                                  167308-34-1P, Iron sodium tungsten
     167308-31-8P
     oxide silicate (Fe5Na5W8O32(SiO4)) 167308-35-2P 167308-36-3P
     167308-37-4P 167308-38-5P 167397-05-9P 167397-06-0P
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     (Biological study, unclassified); SPN (Synthetic preparation); THU
     (Therapeutic use); BIOL (Biological study); PREP (Preparation);
    USES (Uses)
        (polyoxometallates for treatment of flavivirus
        infections)
IT
    140186-99-8
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     (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (polyoxometallates for treatment of flavivirus
        infections)
ΙT
     6834-92-0 7558-80-7, Sodium dihydrogen phosphate
     Cobalt chloride, reactions 7699-43-6
                                            10421-48-4, Ferric
             13472-45-2 64684-57-7
    RL: RCT (Reactant); RACT (Reactant or reagent)
        (polyoxometallates for treatment of flavivirus
        infections)
     63950-53-8P 102073-48-3P 114594-66-0P 139631-90-6P
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                  150923-48-1P 152166-48-8P
                                                167308-25-0P
     167308-26-1P
                  167308-28-3P 167308-33-0P
     RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (polyoxometallates for treatment of flavivirus
        infections)
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L114 ANSWER 45 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

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ACCESSION NUMBER:
                         1995:751176 HCAPLUS
DOCUMENT NUMBER:
                         123:267525
TITLE:
                         Synthetic and catalytic studies of
                         inorganically pillared and organically
                         pillared layered double hydroxides
                         Don Wang, J.; Serrette, Genevieve; Tian, Ying;
AUTHOR (S):
                         Clearfield, Abraham
CORPORATE SOURCE:
                         Department of Chemistry, Texas A and M
                         University, College Station, TX, 77843, USA
                         Applied Clay Science (1995), 10(1-2), 103-15 CODEN: ACLSER; ISSN: 0169-1317
SOURCE:
PUBLISHER:
                         Elsevier
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     By using freshly prepared or thoroughly wet layered double
     hydroxides (LDHs), or by copptg. the LDHs in the presence of the
     pillaring anions, we have synthesized and performed a catalytic
     study on a large number of pillared HT anionic clays. The metal
     cations used in the layers include Al3+, Cr3+, Fe3+, Mg2+, Ni2+,
     Zn2+, Co2+, Fe2+, Cu2+, Ba2+, and Ca2+. Pillaring was achieved
     with an extensive variety of heteropolyoxometalates having the
     Keggin structure, as well as some other larger
     heteropolyoxometalate anions. The product was usually a
     mixture of one phase with an ideally pillared structure and
     another phase containing defects in the layers; defects that were
     created during the pillaring reaction. Hexacyanoferrate(III) was
     also used as a pillaring anion in a clean reaction that
     provides an example of LDHs pillared with organometallic species.
     The organic pillars, [O3P-C6H4-PO3]4-, [O3P-C6H4-C6H4-PO3]4-, and
     [O3P-C6H4-C6H4-C6H4-PO3]4-, first formed intercalates with
     d-spacings exactly equal to the sum of the height of the pillar
     and the thickness of the LDH layer. These intercalates, in turn,
     dehydrate easily to yield lower d-spacing compds. in which the
     organic pillars are, presumably, fixed onto the layers through P-O-M
     bonds. The catalysis tests of polyoxometalate-pillared
     LDHs on isopropanol showed high levels of conversion, generally
     enhanced acid activity, and a gradual change in selectivity with
     respect to dehydrogenation vs. dehydration. The selectivity data
     are presented in correlation with the composition of the LDH
     layers.
     144317-57-7 144435-43-8 169052-67-9
     169052-73-7 169052-74-8
     RL: CAT (Catalyst use); USES (Uses)
        (synthetic and catalytic studies of inorganically pillared and
        organically pillared layered double hydroxides)
RN
     144317-57-7 HCAPLUS
CN
    Vanadate(6-), nona-μ-οxotrioxo(pentadeca-μ-
     oxononaoxononatungstate) [µ12-[phosphato(3-)-
     0:0:0:0':0':0':0'':0'':0'':0''':0''']]tri-, magnesium
     (OC-6-11)-hexahydroxyaluminate(3-) hydroxide (1:30:6:36) (9CI)
     (CA INDEX NAME)
     CM
          1
     CRN 63454-68-2
     CMF 040 P V3 W9
     CCI CCS
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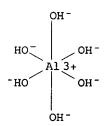
^{*} STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





CM 2

CRN 18893-33-9 CMF Al H6 O6 CCI CCS



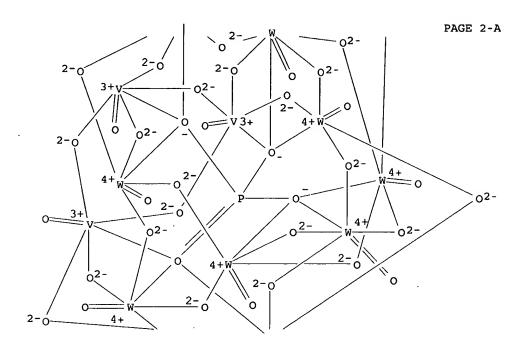
RN 144435-43-8 HCAPLUS

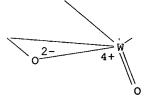
CN Vanadate(6-), nona-µ-oxotrioxo(pentadeca-µ-oxononaoxononatungstate)[µ12-[phosphato(3-)-0:0:0:0':0':0':0':0'':0'':0'':0'']]tri-, magnesium (OC-6-11)-hexahydroxyaluminate(3-) (1:12:6) (9CI) (CA INDEX NAME)

CM 1

CRN 63454-68-2 CMF 040 P V3 W9 CCI CCS

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





PAGE 3-A

CM 2

CRN 18893-33-9 CMF Al H6 O6 CCI CCS

Les Henderson

RN 169052-67-9 HCAPLUS

CN Tungstate(3-), tetracosa-μ-oxododecaoxo[μ12-[phosphato(3-)-0:0:0:0':0':0':0'':0'':0''':0''']]dodeca-, magnesium (OC-6-11)-hexahydroxyaluminate(3-) hydroxide (1:12:3:12) (9CI) (CA INDEX NAME)

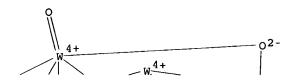
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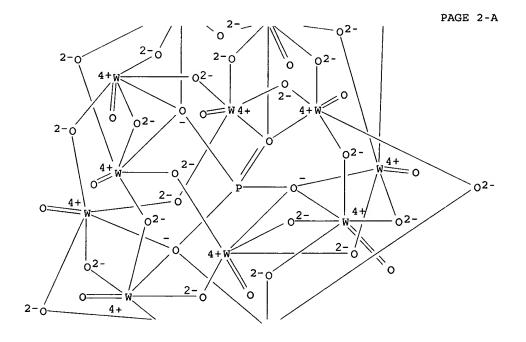
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CM 2

CRN 12534-77-9 CMF 040 P W12 CCI CCS

PAGE 1-A





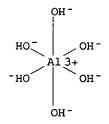
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PAGE 3-A

RN 169052-73-7 HCAPLUS

CM 1

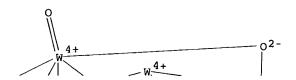
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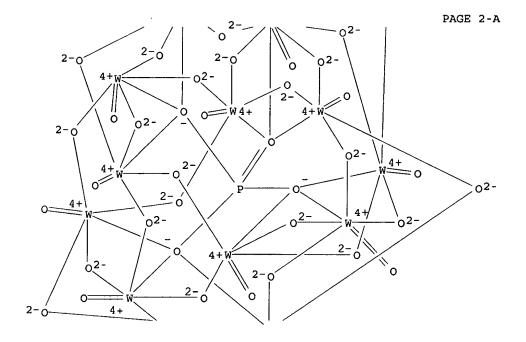


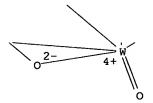
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CRN 12534-77-9 CMF 040 P W12 CCI CCS

PAGE 1-A



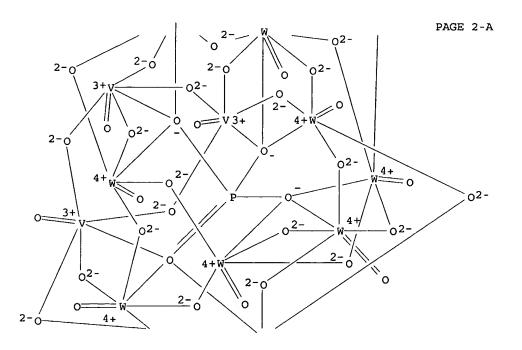




cci ccs

PAGE 3-A

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT



0²⁻ 4+ W

PAGE 3-A

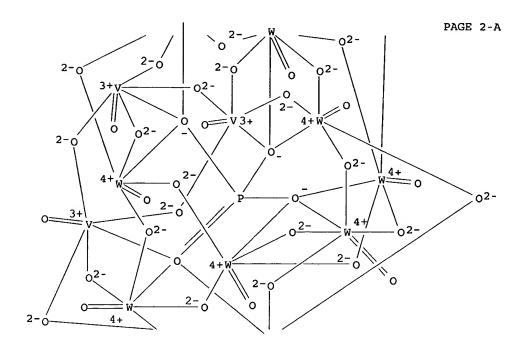
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CRN 18893-33-9
CMF Al H6 O6
CCI CCS

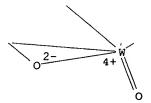
CCI CCS

IT 169052-71-5P 169052-72-6P RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses) (synthetic and catalytic studies of inorganically pillared and organically pillared layered double hydroxides) 169052-71-5 HCAPLUS RNCN Vanadate(6-), nona- μ -oxotrioxo(pentadeca- μ oxononaoxononatungstate) (µ12-[phosphato(3-)-0:0:0:0':0':0':0'':0'':0'':0''':0''']]tri-, nickel(2+) (OC-6-11)-hexahydroxyaluminate(3-) hydroxide (1:30:6:36) (9CI) (CA INDEX NAME) CM 1 CRN 63454-68-2 CMF 040 P V3 W9

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

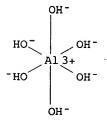
Les Henderson Page 348 571-272-2538





CM

CRN 18893-33-9 CMF Al H6 O6 CCI CCS

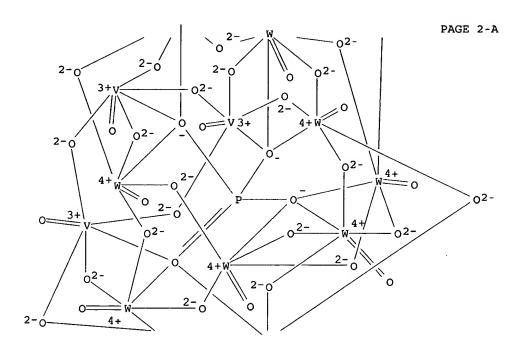


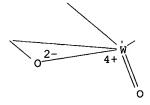
RN

169052-72-6 HCAPLUS Vanadate(6-), nona- μ -oxotrioxo(pentadeca- μ -CN oxononaoxononatungstate) [μ12-[phosphato(3-)0:0:0:0':0':0'':0'':0''':0''':]]tri-, copper(2+)
(OC-6-11)-hexahydroxyaluminate(3-) (1:12:6) (9CI) (CA INDEX NAME) CM

CRN 63454-68-2 CMF 040 P V3 W9 CCI CCS

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





PAGE 3-A

CM 2

CRN 18893-33-9

CMF Al H6 O6

CCI CCS

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OH:
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CC 67-1 (Catalysis, Reaction Kinetics, and Inorganic Reaction Mechanisms)

Section cross-reference(s): 53, 78

69048-27-7, ΙT 13408-62-3, Hexacyanoferrate(III) 69048-26-6 Aluminum magnesium carbonate hydroxide (Al2Mg4(CO3)(OH)12) 144317-57-7 144435-43-8 169052-67-9

169052-68-0 169052-69-1 169052-70-4 169052-73-7 169052-74-8 169112-15-6 169112-16-7 169112-17-8 169112-18-9

RL: CAT (Catalyst use); USES (Uses)

(synthetic and catalytic studies of inorganically pillared and organically pillared layered double hydroxides)

169052-71-5P 169052-72-6P IT

RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(synthetic and catalytic studies of inorganically pillared and organically pillared layered double hydroxides)

L114 ANSWER 46 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1995:574425 HCAPLUS

DOCUMENT NUMBER:

123:47421

TITLE:

Of therapy, toxicity and tungstates:

the anti-retroviral pharmacology of

polyoxometalates

AUTHOR(S):

Blasecki, John W.

CORPORATE SOURCE: Viral Diseases Research, DuPont Merck

Pharmaceutical Company, Glenolden, PA, 19036,

USA

SOURCE:

Topics in Molecular Organization and

Engineering (1994), 10, 373-85 CODEN: TMOEE7; ISSN: 0927-0817

DOCUMENT TYPE:

Journal

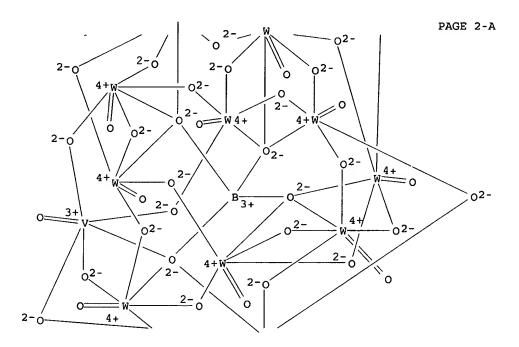
LANGUAGE: English

Based upon activity in a number of in vitro assays of anti-retroviral activity, E3925 (alphavanadoundecatungstoboric acid, hexa-potassium salt (K6[BVW11040] ·xH20)) was elected for further evaluation as a potential candidate for clin. treatment of HIV/AIDS. When tested in mice against Friend leukemia virus (FLV), which was used as a surrogate model of retrovirus-induced immuno-deficiency disease, E3925 was as efficacious as AZT and superior to HPA-23. Treatment of FLV-infected mice could be delayed up to 48 h post-infection with no significant loss of antiviral activity. Careful manipulation of loading dose, maintenance dose and administration interval demonstrated that anti-retroviral efficacy could be maintained by administering E3925 as infrequently as every two weeks. In combination therapy expts., E3925 was both compatible and additive with AZT in reducing the severity of FLV-induced disease in mice. Further clin. development of this compound was prevented by treatment-related coagulopathy.

TΤ 93253-86-2

> RL: ADV (Adverse effect, including toxicity); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study);

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT



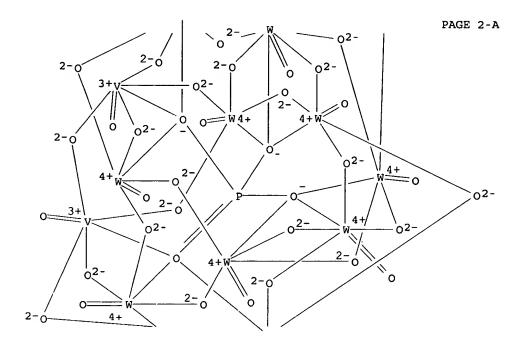


●6 K+

IT 12773-19-2 108834-31-7 140156-49-6
 RL: BAC (Biological activity or effector, except adverse); BSU
 (Biological study, unclassified); THU (Therapeutic use); BIOL
 (Biological study); USES (Uses)
 (anti-retroviral pharmacol. of polyoxometalates)
RN 12773-19-2 HCAPLUS
CN Vanadate(5-), (heptadeca-μ-oxodecaoxodecatungstate)hepta-μ oxodioxo[μ12-[phosphato(3-)-0:0:0':0':0':0':0'':0'':0'':0'']

':O''']]di-, pentapotassium (9CI) (CA INDEX NAME)

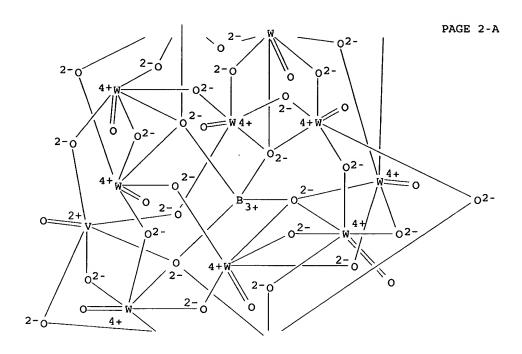
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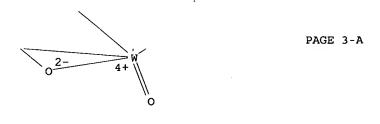




●5 K+

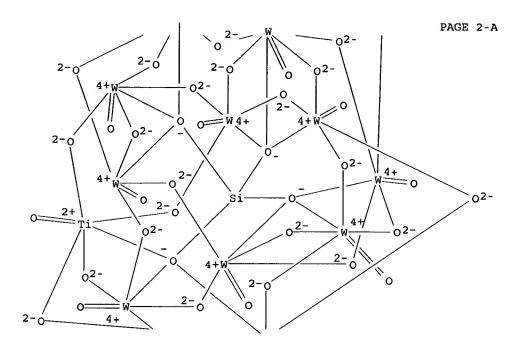
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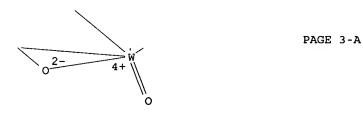




●7 K+

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





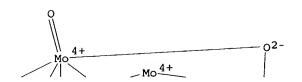
●6 K+

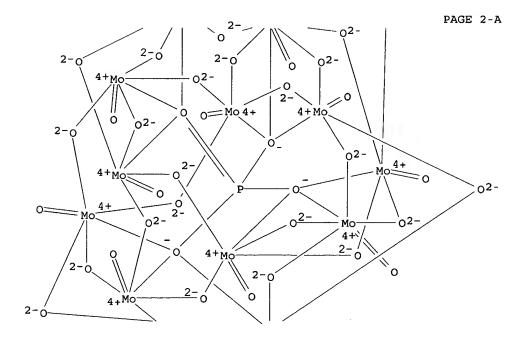
CC

```
1-5 (Pharmacology)
     antiretroviral polyoxometalate E3925
ST
     vanadoundecatungstoborate
ΙT
     Virucides and Virustats
        (anti-retroviral pharmacol. of polyoxometalates)
IT
     Virus, animal
        (retro-, anti-retroviral pharmacol. of polyoxometalates
IT
     93253-86-2
     RL: ADV (Adverse effect, including toxicity); BAC (Biological
     activity or effector, except adverse); BSU (Biological study,
     unclassified); THU (Therapeutic use); BIOL (Biological study);
     USES (Uses)
        (anti-retroviral pharmacol. of polyoxometalates)
ΙT
     12773-19-2 108834-31-7 140156-49-6
     RL: BAC (Biological activity or effector, except adverse); BSU
     (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (anti-retroviral pharmacol. of polyoxometalates)
IT
     30516-87-1, Azt
     RL: BAC (Biological activity or effector, except adverse); BSU
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```
(Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (anti-retroviral pharmacol. of polyoxometalates and)
     101346-86-5
ΙT
     RL: BAC (Biological activity or effector, except adverse); BSU
     (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (santi-retroviral pharmacol. of polyoxometalates)
L114 ANSWER 47 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1994:691255 HCAPLUS
DOCUMENT NUMBER:
                         121:291255
TITLE:
                         polyoxometalates: a class of
                         compounds with remarkable topology
AUTHOR(S):
                         Del Gado, O.; Dress, A.; Mueller, A.; Pope, M.
CORPORATE SOURCE:
                         Dep. Math., Univ. Bielefeld, Bielefeld,
                         D-33501, Germany
SOURCE:
                         Molecular Engineering (1993), 3(1-3), 9-28
                         CODEN: MOLEEV; ISSN: 0925-5125
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     Structures of polyoxometalates frequently are discovered
     to be based upon regular convex polyhedra, including the Platonic
     and Archimedean solids. A topol. approach involving barycentric
     subdivision of the faces of such polyhedra, leads to their
     description as combinations of triangular building
    blocks assembled according to systematic rules. An anal. of the
    Keggin structure of [Mo12036(PO4)]3-, is presented. It is the
     only spherical polyhedral structure of T symmetry built up from 6
     8-gons and 8 6-gons. Similarly, there is only 1 spherical
     polyhedral structure of T symmetry built up from 8 6-gons and
     twenty-4 4-gons satisfying also some obvious chemical
     combinatorial constraints. Such a structure is observed for
     [H9V18O42(VO4)]6-. Anal. of possible structures of lower symmetry
     (D3, D4), e.g. as observed for [V15O36(Hal)]6- and [H4V18O42(Hal)]9-,
     reveals the onset of combinatorial explosion.
     For example, there are 67 D3-structures satisfying the chemical
    condition.
IT
    12379-13-4, Molybdophosphate (Mol2036(PO4)3-)
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (topol. of)
RN
    12379-13-4 HCAPLUS
    Molybdate(3-), tetracosa-\mu-oxododecaoxo[\mu12-[phosphato(3-)-
CN
    0:0:0:0':0':0':0'':0'':0'':0''':0''']]dodeca- (9CI) (CA
    INDEX NAME)
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PAGE 1-A





02- Mo4+

78-7 (Inorganic Chemicals and Reactions)

PAGE 3-A

```
ST
     polyoxometalate classification topol; barycentric
     subdivision polyoxometalate; combinatorial
     explosion polyoxometalate; Keggin
     polyoxometalate topol
     Molecular topology
ΙT
        (of polyoxometalates)
     12379-13-4, Molybdophosphate (Mo12036(PO4)3-)
тт
     158868-03-2
                   158868-04-3, Vanadate (V170415-)
                                                       158908-66-8
     158908-68-0
                   158916-13-3
                                 158916-14-4
                                               158925-41-8
     158925-53-2
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (topol. of)
L114 ANSWER 48 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1994:664655 HCAPLUS
DOCUMENT NUMBER:
                         121:264655
TITLE:
                         A More General Approach to Distinguishing
                         "Homogeneous" from "Heterogeneous" Catalysis:
                         Discovery of Polyoxoanion- and
                         Bu4N+-Stabilized, Isolable and Redissolvable,
                         High-Reactivity Ir.apprx.190-450 Nanocluster
                         Catalysts
AUTHOR(S):
                         Lin, Yin; Finke, Richard G.
CORPORATE SOURCE:
                         Department of Chemistry, Colorado State
                         University, Ft. Collins, CO, 80523, USA
SOURCE:
                         Inorganic Chemistry (1994), 33(22), 4891-910
                         CODEN: INOCAJ; ISSN: 0020-1669
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     A more general approach to distinguishing between so-called
     homogeneous vs. heterogeneous catalysts has been developed and
     intrinsically tested in answering the question "what is the true
     catalyst in the active hydrogenation system which evolves from
     cyclohexene, hydrogen, and the discrete, polyoxoanion-supported
     Ir(I) catalyst precursor (Bu4N)5Na3[(1,5-
     COD) Ir · P2W15Nb3O62] ". The approach developed and utilized
     consists of four categories of expts.: (i) catalyst isolation and
     characterization studies, with an emphasis initially on TEM; (ii)
     initial kinetic studies, emphasizing whether or not the isolated
     catalyst can account for the observed kinetics, especially any induction
     period seen, and whether or not the reaction exhibits a ±10%
     reproducible rate; (iii) quant. phenomenol. catalyst poisoning and recovery expts.; (i.v.) addnl. kinetic and
     mechanistic studies and chemical tests, all interpreted with strict
     adherence to the principle that the correct description of the
     catalyst (i.e., the correct mechanism) will explain all of the
     data. The present approach has identified a previously unknown
     type of hybrid homogeneous-heterogeneous, Ir.apprx.190-
     450 · polyoxoanion/Bu4N+ catalyst of average composition
     [Ir(0).apprx.300(P4W30Nb6O12316-).apprx.33](Bu4N).apprx.300Na.appr
     x.233. A min. mechanistic scheme for the catalyst's evolution,
```

CC

consisting of the autocatalytic generation of the Ir.apprx.190-450 nanoclusters, is shown to account for all of the observed results, including the findings of the rate-enhancing effects of H+, H2O,

and acetone impurities that were puzzling in the earlier stages of this work.

IT 92762-46-4P

RL: CAT (Catalyst use); PNU (Preparation, unclassified); PREP (Preparation); USES (Uses)

(preparation and characterization of polyoxoanion- and tetrabutylammonium-stabilized iridium nanocluster hydrogenation catalysts)

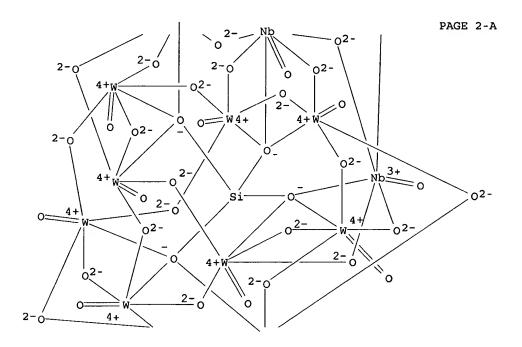
RN 92762-46-4 HCAPLUS

Niobate(7-), [μ 12-[orthosilicato(4-)- κ 0: κ 0: κ 0': κ 0': κ 0': κ 0' CN

':k0'':k0''':k0''':k0''']]nona-

 μ -oxotrioxo(pentadeca- μ -oxononaoxononatungstate)tri- (9CI) (CA INDEX NAME)

STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





IT 153299-14-0

RL: CAT (Catalyst use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)

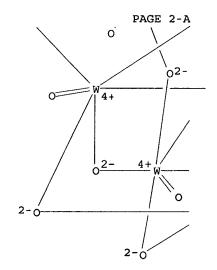
(preparation and characterization of polyoxoanion- and tetrabutylammonium-stabilized iridium nanocluster hydrogenation catalysts)
RN 153299-14-0 HCAPLUS
CN 1-Butanaminium, N,N,N-tributyl-, sodium [[(1,2,5,6-η)-1,5-cyclooctadiene]iridate] [μ12-[orthosilicato(4-)-κ0:κ0:κ0:κ0':κ0':κ0'':κ0''']] hepta-μ-οxotri-μ3-oxotrioxo[tetradeca-μ-οxononaoxononatungstate]triniobate(6-) (4:2:1) (9CI) (CA INDEX NAME)

CM 1

CRN 153299-12-8

CMF C8 H12 Ir Nb3 O40 Si W9
CCI CCS

- * STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY AVAILABLE VIA OFFLINE PRINT
- * STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY AVAILABLE VIA OFFLINE PRINT



* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

CM 2

CRN 10549-76-5 CMF C16 H36 N

CC 67-1 (Catalysis, Reaction Kinetics, and Inorganic Reaction

```
Mechanisms)
ΙT
     114691-25-7P
        catalysts)
IT
ACCESSION NUMBER:
DOCUMENT NUMBER:
TITLE:
INVENTOR(S):
```

Section cross-reference(s): 22, 78

7439-88-5DP, Iridium, nanoclusters 92762-46-4P

RL: CAT (Catalyst use); PNU (Preparation, unclassified); PREP (Preparation); USES (Uses)

(preparation and characterization of polyoxoanion- and tetrabutylammonium-stabilized iridium nanocluster hydrogenation

152075-49-5 153299-14-0

RL: CAT (Catalyst use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)

(preparation and characterization of polyoxoanion- and tetrabutylammonium-stabilized iridium nanocluster hydrogenation catalysts)

L114 ANSWER 49 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

1994:525210 HCAPLUS

121:125210

Heteropolytungstates as antiviral agents Weigold, Helmut; Bartholomeusz, Angeline Ingrid; Marcuccio, Sebastian Mario; Holan,

ADDITION NO

חאיים

George

PATENT ASSIGNEE(S):

Commonwealth Scientific and Industrial

Research Organization, Australia

SOURCE:

PCT Int. Appl., 32 pp. CODEN: PIXXD2

שתעת

DOCUMENT TYPE:

Patent

LANGUAGE:

English

KIMD

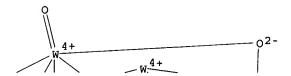
FAMILY ACC. NUM. COUNT:

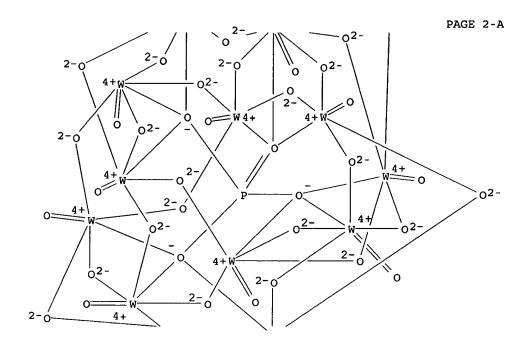
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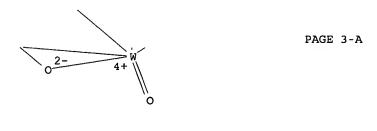
PF	PATENT NO.						KIND DATE			APPLICATION NO.						DATE	
							•										
WC)	9412192				A1		19940609		WO 1993-AU606						1993	
														1129			
		W:	AT,	AU,	BB,	BG,	BR,	BY,	CA,	CH,	CZ,	DE,	DK,	ES,	FI,		
			-	_	-						MG, US,			NL,	NO,	NZ,	
		RW:	-	-		•			•	•	GR,			LU,	MC,	NL,	
			PT,	SE,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	ML,	MR,	NE,	SN,	
	_		TD,											_			
AU 9455545					Al	A1 19940622 AU 1994-55545							5				
																1993	
GT 1002006								OV 1002 101510						1129			
CN 1093906					A 19941026				CN 1993-121712						1000		
																1993 1201	
PRIORIT	v	ΔΡΡΙ	ר זא.	NFO							AU 19	992-6	5115		1	1201	
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																1201	
									1	WO 19	993-1	AU606	5	Ţ	1		
																1993	
																1129	

AB Heteropolytungstates having antiflaviviral activity has the general formula (I) to (VII): (I) An[XW12040] wherein X is selected from PV, SiIV, GeIV, CoII, CoIII, ZnII, CuII, BIII, HI, AlIII, FeIII, VV, GaIII, MnIV, CIV; (II) An[X2W18062] wherein X is PV; (III) An[XW11039] wherein X is selected from PV, SiIV, GeIV, BIII, AlIII, GaIII, FeIII, CoIII; (IV) An[XW9O34] wherein X is selected from PV, SiIV, GeIV; (V) An[X2W17-mMmO61] wherein X is PV, M is MoVI, and m is 0, 1, 2, 4, or 5; (VI) An[X2W15-mMmO56]

wherein X is PV, M is VV or MoVI, and m is 0 or 1; (VII) An [XM2W9O39] wherein X is P, and M is Zr; and wherein in each of the general formula (I) to (VII), A is a cation, and n is the number of cations necessary for elec. neutrality of the mol.; or dimers, hydrates of pharmaceutically acceptable derivs. thereof. Pharmaceutical compns. and methods for the treatment or prophylaxis of a flaviviral-associated infection which involve the use of these compds. are also disclosed. For example, 23 compds. including H3PW1240 nH2O were tested for their ability to inhibit RNA synthesis in an in vitro polymerase assay. 12266-04-5 12357-89-0 12501-23-4 IT 56127-18-5 86692-07-1 133515-28-3 134879-28-0 157208-01-0 RL: BIOL (Biological study) (flavivirus-associated infections treatment with) RN 12266-04-5 HCAPLUS Tungstate(3-), tetracosa- μ -oxododecaoxo[μ 12-[phosphato(3-)-CN κ0:κ0:κ0:κ0':κ0':κ0':κ0' ':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec a-, trisodium, hydrate (9CI) (CA INDEX NAME)



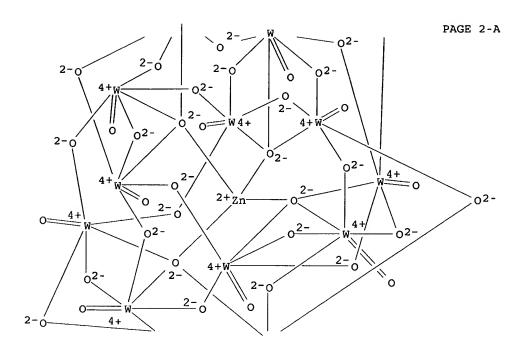


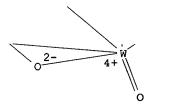


●3 Na+

●x H₂O

RN 12357-89-0 HCAPLUS
CN Tungstate(6-), tetracosa-μ-oxotetra-μ4 oxododecaoxozincatedodeca-, hexapotassium, hydrate (9CI) (CA
 INDEX NAME)





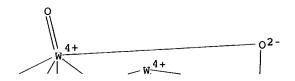
●6 K+

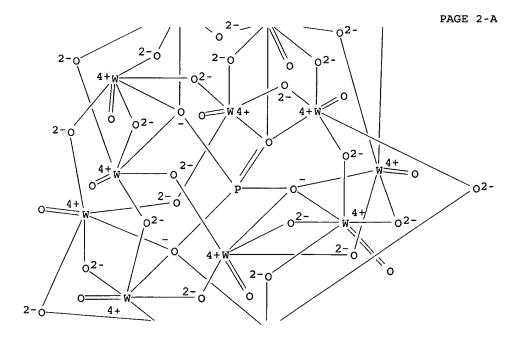
●x H₂O

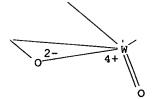
RN 12501-23-4 HCAPLUS CN

Tungstate(3-), tetracosa-µ-oxododecaoxo[µ12-[phosphato(3-)-0:0:0':0':0':0'':0'':0''':0''']]dodeca-, trihydrogen, hydrate (9CI) (CA INDEX NAME)

PAGE 1-A



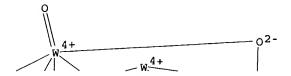


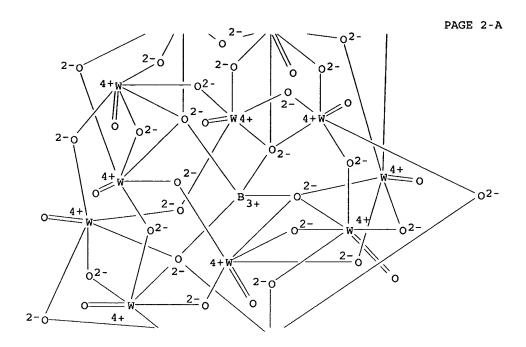


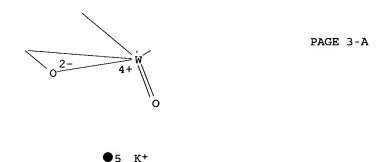
●3 H+

●x H₂O

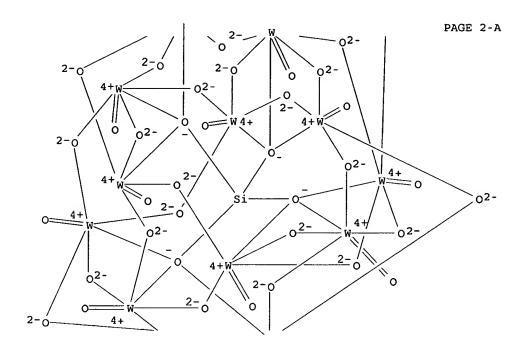
RN 56127-18-5 HCAPLUS
CN Tungstate(5-), tetracosa-μ-oxododecaoxo[μ12[tetrahydroxyborato(5-)-κ0:κ0:κ0:κ0':.kapp
a.O':κ0'':κ0'':κ0'':κ0'':.kappa
.O''':κ0''']]dodeca-, pentapotassium, hydrate (9CI) (CA INDEX NAME)

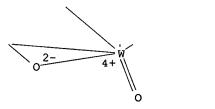






●x H₂O





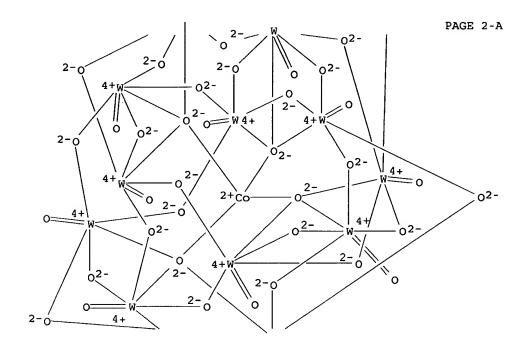
•4 Na+

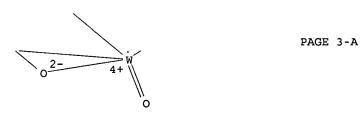
●x H₂O

Tungstate(6-), cobaltatetetracosa- μ -oxotetra- μ 4-

CN

oxododecaoxododeca-, hexapotassium, hydrate (9CI) (CA INDEX NAME)





●6 K+

●x H₂O

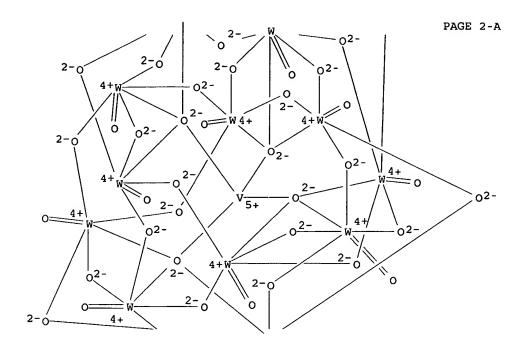
RN 157208-01-0 HCAPLUS
CN Vanadate(3-), tetra-µ4-oxo(tetracosa-µoxododecaoxododecatungstate)-, trihydrogen, compd. with
N,N-dimethylmethanamine (1:3), hydrate (9CI) (CA INDEX NAME)

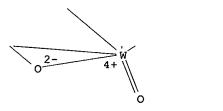
CM 1

CRN 157208-00-9

CMF H . 1/3 O40 V W12

CCI CCS





●3 H+

CM 2

CRN 75-50-3 CMF C3 H9 N

CH₃ H₃C-N-CH₃

IC

ICM A61K033-24 ICS C01G041-00

CC 1-5 (Pharmacology) IT

12266-04-5 12357-89-0 12501-23-4

56127-18-5 58916-01-1 62682-77-3 **86692-07-1** 114714-81-7 116231-28-8 121796-02-9 113471-17-3

133515-28-3 134879-28-0 136171-80-7

157079-66-8 157111-20-1 157111-21-2 157111-22-3

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157111-23-4 157111-24-5
                           157177-66-7 157208-01-0
RL: BIOL (Biological study)
   (flavivirus-associated infections treatment with)
```

L114 ANSWER 50 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 1994:226955 HCAPLUS

DOCUMENT NUMBER:

120:226955

TITLE:

Pharmaceutical compositions

containing heteropolytungstates for the

treatment of flavivirus infections INVENTOR(S):

Weigold, Helmut; Bartholomeusz, Angeline Ingrid; Holan, George; Marcuccio, Sebastian

Mario; Wright, Peter James

PATENT ASSIGNEE(S):

Commonwealth Scientific and Industrial

Research Organization, Australia

SOURCE:

PCT Int. Appl., 29 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA'	KIND DAT		DATE		API	APPLICATION NO.											
WO	9321934				A1		19931111		WO 1993-AU192					1993 0430			
		MN,	MW,	NO,	NZ,	PL,	RO,	RU,	HU, JI SD, SI GB, GI	C, UA,	US,	VN	-	•			
AU	9342	TD,	TG	·		•			CM, GA		•	·	NE,	·			
HU 71677				A2	A2 19960129 HU 19												
CN 1082892					A	:	1994	0302	CN	1993-		1993 0430					
PRIORITY	APPI	LN.	INFO	. :					AU	1992-	2213		1	1993 0501 A			
									•••					1992 0501			
									WO	1993-	AU192	2	1	1993 0430			

AB Pharmaceutical compns. containing heteropolytungstates and derivs. thereof are used for the treatment or prophylaxis of flavivirus infections, e.g. yellow fever. The inhibitory concentration of K4[C5H5TiPW11039].nH20 (I) against kunjin virus-infected vero cells was at 5-100 µM. A tablet contained I 250, lactose 210, povidone 15, Na starch glycollate 20, and Mg stearate 5mg.

84303-06-0P 93222-18-5P 101144-77-8P

105785-76-0P 152313-54-7P 152313-55-8P

152313-57-0P 152313-58-1P 152313-59-2P

152344-57-5P 152344-58-6P 152344-59-7P

152981-11-8P 154165-09-0P 154165-10-3P

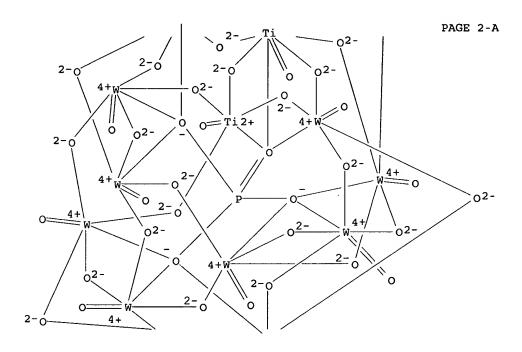
154165-11-4P 154165-13-6P 154165-14-7P

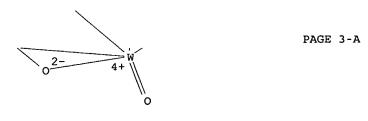
RL: PREP (Preparation)

(preparation of, pharmaceutical compns. containing, for treatment of flavivirus infections)

RN 84303-06-0 HCAPLUS CN Titanate(7-), (heptadeca-μ-oxodecaoxodecatungstate)hepta-μoxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ
':κΟ':κΟ':κΟ'':κΟ'':κΟ''
:κΟ''':κΟ''']]di-, heptapotassium (9CI) (CA INDEX
NAME)

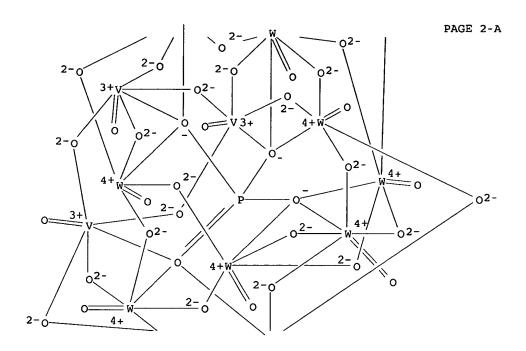
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





●7 K+

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

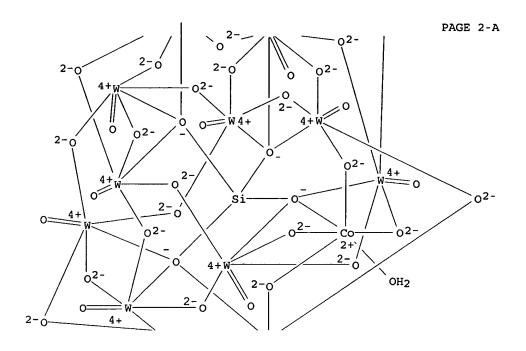


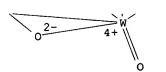


●6 Cs+

```
RN
     101144-77-8 HCAPLUS
CN
     Vanadate(9-), [heptacosa-\mu-oxopentadecaoxo[\mu9-[phosphato(3-)-
     κ0:κ0:κ0:κ0':κ0':κ0'',κ0
    '':κΟ''':κΟ''']]pentadecatungstate]nona-μ-
     oxotrioxo[μ9-[phosphato(3-)-κ0:κ0:κ0:κ0
':κ0':κ0'':κ0'':κ0''':κ0''']]tri-,
     octapotassium hydrogen (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN
    105785-76-0 HCAPLUS
CN
     Tungstate(6-), (aquacobaltate)[\mu12-[orthosilicato(4-)-
     κ0:κ0:κ0:κ0':κ0':κ0':κ0'
     ':κ0'':κ0'':κ0''':κ0''']]tetra
cosa-μ-oxoundecaoxoundeca-, hexapotassium (9CI) (CA INDEX
     NAME)
```

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





PAGE 3-A

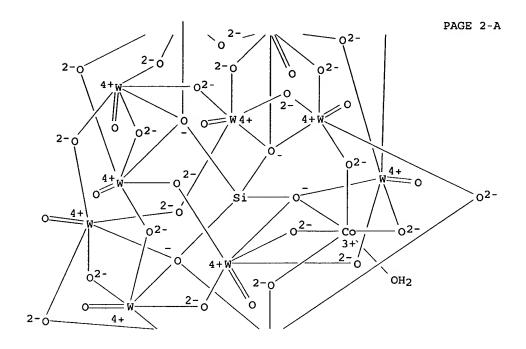
●6 K+

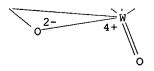
CM 1

CRN 66304-66-3 CMF CO H2 O40 Si W11 . 5 H CCI CCS

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

Les Henderson Page 374 571-272-2538





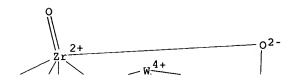
●5 H+

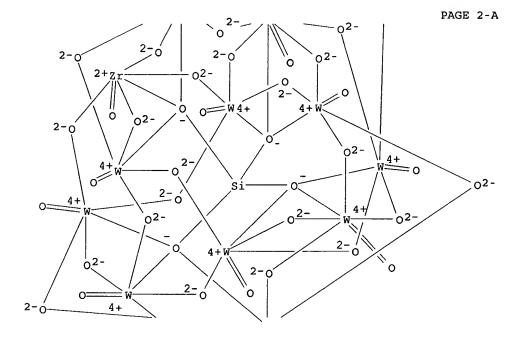
CM

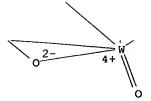
CRN 113-00-8 CMF C H5 N3

RN

152313-55-8 HCAPLUS
Zirconate(8-), (heptadeca-µ-oxodecaoxodecatungstate)[µ12[orthosilicato(4-)-0:0:0':0':0':0'':0'':0''':0''']]hept
a-µ-oxodioxodi-, tripotassium pentahydrogen (9CI) (CA INDEX CN NAME)







●5 H+

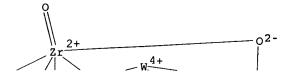
●3 K+

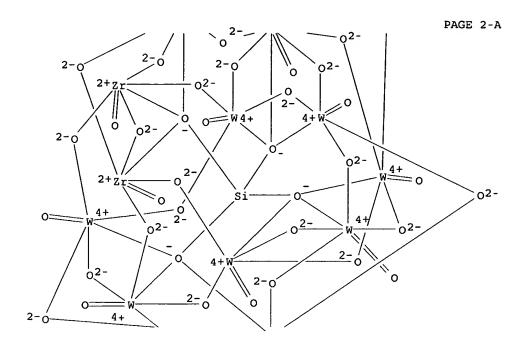
RN 152313-57-0 HCAPLUS

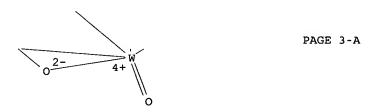
CN Methanaminium, N,N,N-trimethyl-, hydrogen [μ12-[orthosilicato(4-)-0:0:0:0':0':0'':0'':0'':0''':0''']]nona-μoxotrioxo(pentadeca-μ-oxononaoxononatungstate)trizirconate(10-)
(3:7:1) (9CI) (CA INDEX NAME)

CM 1

CRN 152313-56-9 CMF 040 Si W9 Zr3 CCI CCS







CM 2

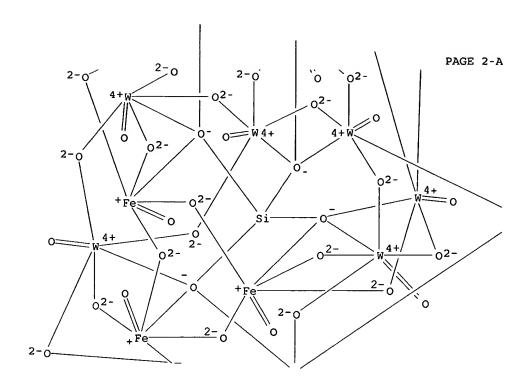
CRN 51-92-3 CMF C4 H12 N

RN 152313-58-1 HCAPLUS

Tungstate(13-), [μ 12-[orthosilicato(4-)- κ 0: κ 0: κ 0: κ 0': κ 0': κ 0': κ 0' CN

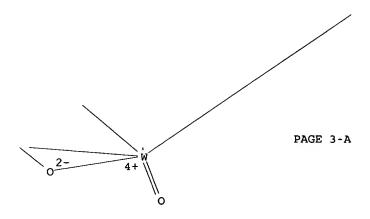
':κ0'':κ0'':κ0''':κ0''':κ0''']]henei

 $cosa-\mu-oxononaoxo(tri-\mu-oxotrioxotriferrate)nona-,$ hexapotassium heptahydrogen (9CI) (CA INDEX NAME)



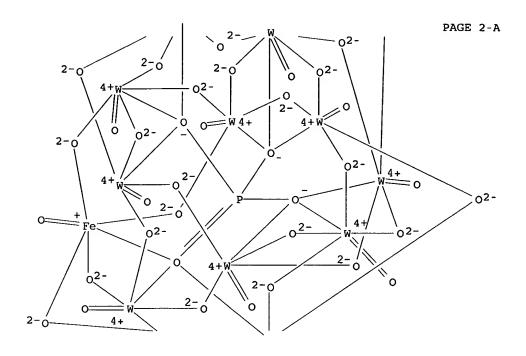
PAGE 2-B

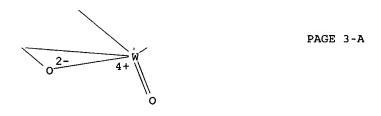
)₀₂-



●7 H+

●6 K+



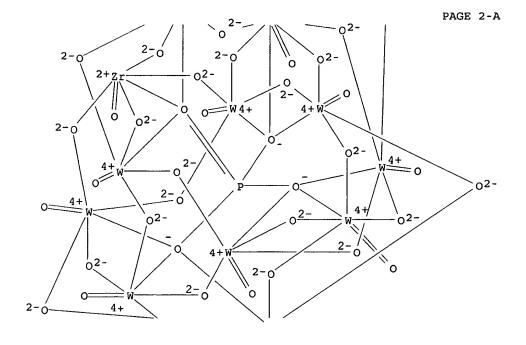


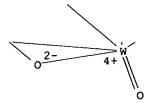
●6 K⁺

RN 152344-58-6 HCAPLUS
CN Zirconate(7-), (heptadeca-μ-oxodecaoxodecatungstate)hepta-μoxodioxo[μ12-[phosphato(3-)-0:0:0':0':0'':0'':0'':0''
':0''']]di-, heptapotassium (9CI) (CA INDEX NAME)

PAGE 1-A

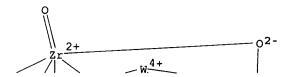


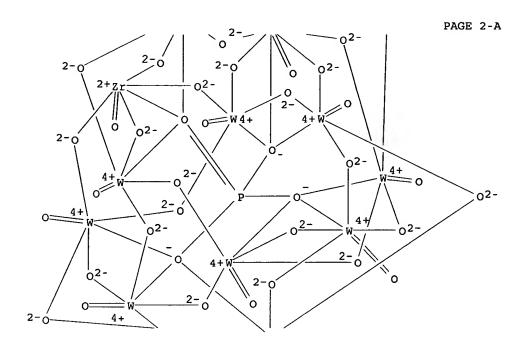


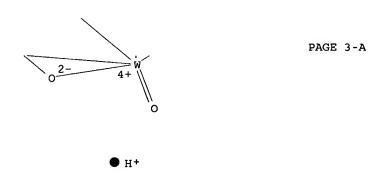


●7 K

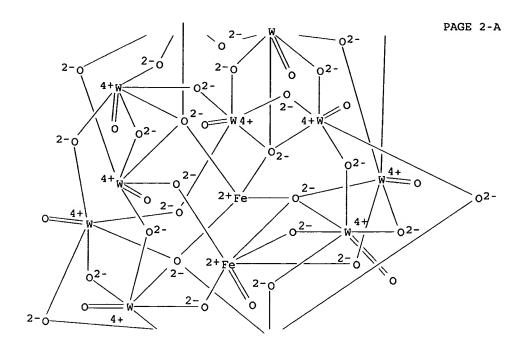
RN 152344-59-7 HCAPLUS
CN Zirconate(7-), (heptadeca-μ-oxodecaoxodecatungstate)hepta-μoxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ
':κΟ':κΟ':κΟ'':κΟ'':κΟ'''
:κΟ''':κΟ''']]di-, hexapotassium hydrogen (9CI) (CA
INDEX NAME)

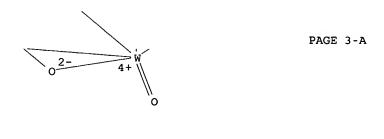






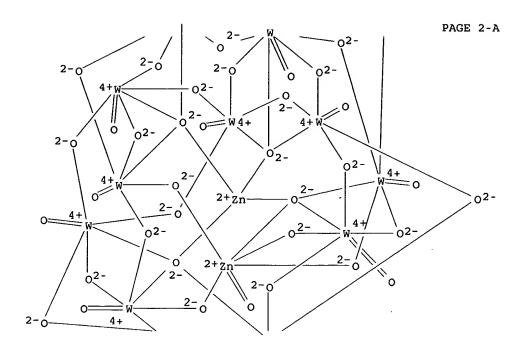
●6 K+

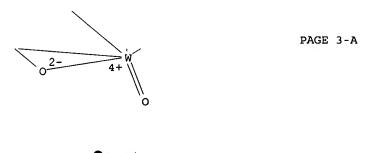




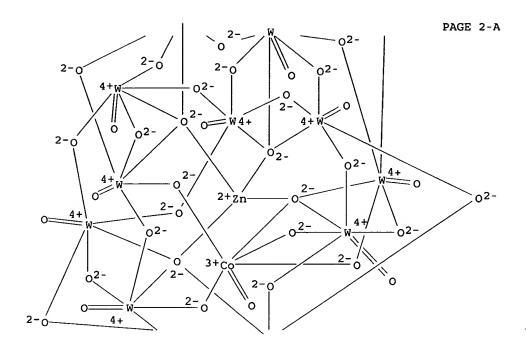
●8 K+

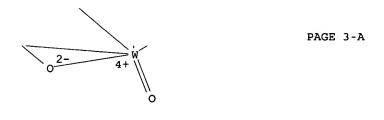
RN 154165-09-0 HCAPLUS
CN Tungstate(8-), tetracosa-μ-oxotetra-μ4oxoundecaoxo(oxodizincate)undeca-, octapotassium (9CI) (CA INDEX NAME)





●8 K+

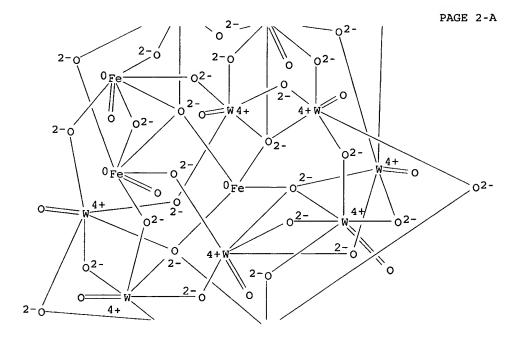




●7 K+

RN 154165-11-4 HCAPLUS CN Tungstate(14-), (di- μ -oxo- μ 4-oxotrioxotetraferrate)docosa- μ -oxotri- μ 4-oxononaoxonona-, hexapotassium octahydrogen (9CI) (CA INDEX NAME)





2- 4+ W

PAGE 3-A

●8 H+

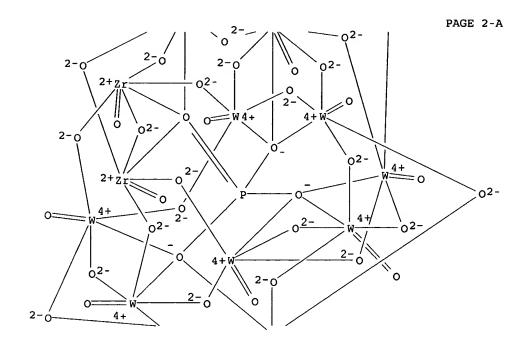
●6 K+

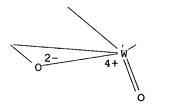
RN 154165-13-6 HCAPLUS
CN Zirconate(9-), nona-µ-oxotrioxo(pentadeca-µoxononaoxononatungstate)[µ12-[phosphato(3-)O:0:0:0':0':0'':0'':0'':0''':0''']]tri-, nonahydrogen,
compd. with N,N-dimethylmethanamine (1:3) (9CI) (CA INDEX NAME)

CM 1

CRN 154165-12-5
CMF H . 1/9 O40 P W9 Zr3
CCI CCS





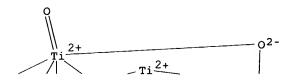


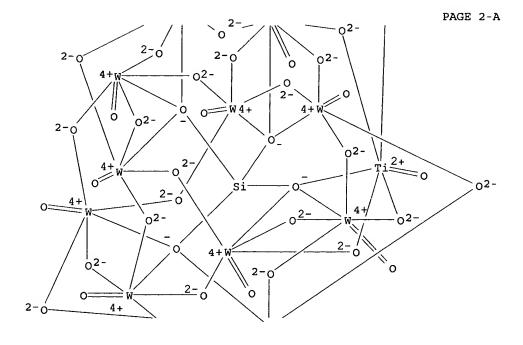
●9 H+

CM 2

CRN 75-50-3 CMF C3 H9 N

PAGE 1-A





02- W

PAGE 3-A

●4 H⁻¹

●6 K+

```
IC
     ICM A61K033-24
     63-6 (Pharmaceuticals)
CC
     Section cross-reference(s): 1, 78
ST
     heteropolytungstate flavivirus infection pharmaceutical
     compn; tablet heteropolytungstate kunjin virus infection;
     yellow fever treatment heteropolytungstate tablet
ΙT
     Virus, animal
        (flavi-, infection with, treatment of, with pharmaceutical
        compns. containing heteropolytungstates)
     Heteropoly acids
IT
     RL: PREP (Preparation)
        (tungstates, preparation of, pharmaceutical compns. containing, for treatment of flavivirus infections)
     63043-34-5P 63043-36-7P 81552-61-6P 81553-24-4P
     84303-06-0P 93222-18-5P 98735-23-0P
                   101347-00-6P
     101144-77-8P
                                  101347-05-1P
                    110717-64-1P
     105785-76-0P
                                   110717-65-2P
                                                  110717-70-9P
     133289-60-8P, Iron sodium tungsten zinc oxide (Fe2Na12W19Zn3O69)
     139901-87-4P, Iron sodium tungsten oxide phosphate
     (Fe2Na7W9O32(PO4))
                         139901-88-5P, Iron potassium tungsten oxide
     phosphate (Fe2K7W9O32(PO4))
                                   140186-99-8P
                                                  141532-38-9P
     141532-43-6P
                   141532-46-9P
                                   141532-49-2P
                                                   141532-61-8P
     141532-65-2P
                    141532-70-9P
                                   141532-71-0P
                                                   147230-49-7P
     152270-86-5P, Cobalt sodium tungsten oxide (Co5Na12W19O68)
     152270-87-6P 152313-54-7P 152313-55-8P
     152313-57-0P 152313-58-1P 152313-59-2P
     152344-57-5P 152344-58-6P 152344-59-7P
     152369-85-2P
                                  152369-87-4P 152444-38-7P
                   152369-86-3P
                                  152444-41-2P 152981-11-8P
     152444-39-8P
                   152444-40-1P
     152992-15-9P, Ammonium iron tungsten oxide phosphate
     ((NH4)7Fe2W9O32(PO4)) 154165-09-0P 154165-10-3P
     154165-11-4P 154165-13-6P 154165-14-7P
     188746-62-5P, Sodium tungsten zinc oxide (Na12W19Zn5O68)
     RL: PREP (Preparation)
        (preparation of, pharmaceutical compns. containing, for
        treatment of flavivirus infections)
L114 ANSWER 51 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1993:93692 HCAPLUS
DOCUMENT NUMBER:
                         118:93692
TITLE:
                         Coated-wire electrodes sensitive to a local
                         anesthetic bupivacaine cation and in vivo
                         applications
AUTHOR (S):
                         Sakate, Hiromu; Wakatsuki, Miho; Kaneshina,
                         Shoji; Yokono, Atsuko; Yokono, Satoshi; Oguli,
                         Kenji
```

CORPORATE SOURCE:

Inst. Coop. Res., Univ. Tokushima, Tokushima,

770, Japan

SOURCE:

Bunseki Kagaku (1992), 41(11), 573-80 CODEN: BNSKAK; ISSN: 0525-1931

DOCUMENT TYPE:

Japanese

Journal LANGUAGE:

A coated-wire electrode sensitive to the local anesthetic bupivacaine cation was prepared by coating copper wire (0.6 or 0.2 mm diameter) with a polyvinylchloride (PVC) membrane of 0.3-0.6 mm thickness. The PVC membrane included a plasticizer and an ion exchanger forming an ion-pair with bupivacaine. Five ion exchangers and 3 plasticizers were used for the electrode membrane. The response times and the slopes of the electromotive force vs. bupivacaine concentration curves were examined by electrodes with various membrane compns. The best results were obtained with the electrode with a PVC membrane composition as follows: (1) ion-exchangers sodium tetrakis[3,5-bis(trifluoromethyl)phenyl]bora te (1-5 mg) or dodecatungstophosphoric acid (2-5 mg) or ion-pair bupivacaine-dodecatungstophosphate (2 mg); (2) plasticizers dioctyladipate (100 mg), dioctylsebacate (100 mg), or dioctylphthalate (150 mg); (3) PVC (100 mg); and (4) solvent THF $(1.5 \ \text{mL})$. The electrodes showed linear responses with Nernstian slopes (58-63 mV/decade) over a concentration range of 3 + 10-5-10-2 M. In measurements with the electrode incorporating dodecatungstophosphoric acid or its ion-pair with bupivacaine used as the sensing material, the change in the pH within a range of 2-8 did not affect the electrode potential at 10-4 M bupivacaine. None of the inorg. and organic cations in biol. samples interfered, as judged by the very small values of the selectivity coeffs. selectivity coeffs. of the bupivacaine electrode toward other local anesthetics decreased in the order of dibucaine > tetracaine > mepivacaine = lidocaine = procaine. A bupivacaine electrode with the PVC membrane, in which bupivacaine-dodecatungstophosphate ion-pair and dioctylphthalate were incorporated, was applied to monitor bupivacaine concns. in the arterial blood of rabbits. The electrode is suitable for measurements of bupivacaine concns. in vivo.

TT 1343-93-7 146090-83-7

RL: ANST (Analytical study)

(in electrode selective for bupivacaine determination in blood)

RN 1343-93-7 HCAPLUS

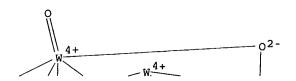
CN Tungstate (3-), tetracosa- μ -oxododecaoxo [μ 12-[phosphato(3-)-

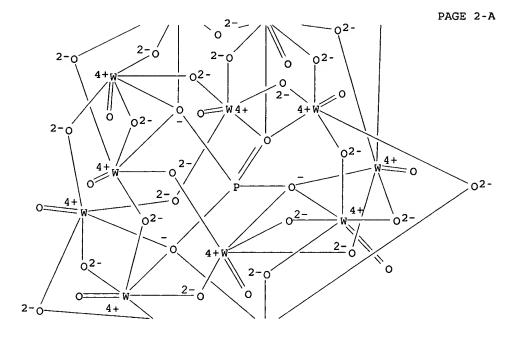
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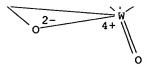
':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec

a-, trihydrogen (9CI) (CA INDEX NAME)

PAGE 1-A







●3 H+

146090-83-7 HCAPLUS RN

Tungstate(3-), tetracosa-µ-oxododecaoxo[µ12-[phosphato(3-)-0:0:0':0':0'':0'':0'':0''':0''']]dodeca-, trihydrogen, compd. with 1-butyl-N-(2,6-dimethylphenyl)-2-piperidinecarboxamide (1:1) (9CI) (CA INDEX NAME)

CM

CN

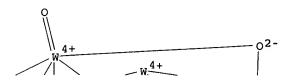
CRN 38396-39-3 CMF C18 H28 N2 O

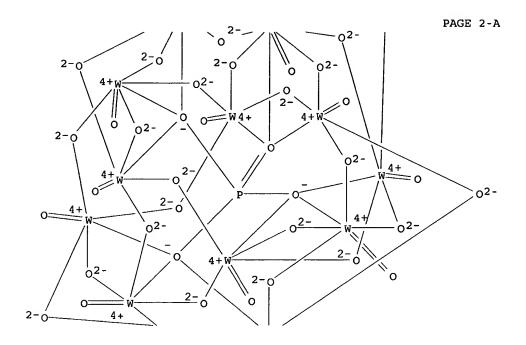
2 CM

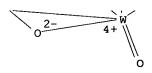
CRN 1343-93-7 CMF H . 1/3 O40 P W12

CCI CCS

PAGE 1-A







FAMILY ACC. NUM. COUNT:

PAGE 3-A

●3 H+

```
CC
      1-1 (Pharmacology)
      Section cross-reference(s): 72
     103-23-1, Dioctyl adipate 117-81-7, Dioctyl phthalate 122-62-3, Dioctyl sebacate 1343-93-7 9002-86-2, Polyvinylchloride 79060-88-1, Sodium tetrakis[3,5-
IT
     bis(trifluoromethyl)phenylborate 146090-83-7
      RL: ANST (Analytical study)
         (in electrode selective for bupivacaine determination in blood)
L114 ANSWER 52 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                             1992:638711 HCAPLUS
DOCUMENT NUMBER:
                             117:238711
TITLE:
                             Kandite clay compositions, and their
                             manufacture
INVENTOR(S):
                             Vaughan, David Evan William
PATENT ASSIGNEE(S):
                             Exxon Research and Engineering Co., USA
                             Eur. Pat. Appl., 17 pp.
SOURCE:
                             CODEN: EPXXDW
DOCUMENT TYPE:
                             Patent
LANGUAGE:
                             English
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PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 491520	A1	19920624	EP 1991-311556	
BF 451520	VI	19920024	EF 1991-311330	1991
				1212
EP 491520	B1	19941026		
R: BE, DE, FR,	GB, IT	, NL		
CA 2055365	AA	19920618	CA 1991-2055365	
				1991
				1113
BR 9105426	Α	19920825	BR 1991-5426	
				1991
JP 05139719	2.2	10020600	TD 1001 331010	1213
JP 05139/19	A2	19930608	JP 1991-331019	1991
				1216
JP 05310414	A2	19931122	JP 1991-361046	1210
31 33323121		20001122	01 1331 301010	1991
				1217
US 5326734	Α	19940705	US 1992-985399	
				1992
				1204
PRIORITY APPLN. INFO.:			US 1990-628514	A
				1990
·				1217
			US 1992-857032	B1
			05 1992-65/032	1992
				0324

AB Pillared interlayered kandite compns. comprise kandite layers separated by inorg. metal or metal oxide pillars derived from charged or neutral oxide, hydroxide, or organometallic clusters containing ≥4 metal atoms. The interlayer distances are substantially greater than those in precursor of the same but nonsepd. clay. The kandite clay compns. are manufactured by (a) intimately mixing finely divided kandite clay with an aqueous solution to obtain a precursor suspension, (b) adding an effective amount of pillaring medium to the dispersion and aging the mixture for a sufficient time to allow infusion of the pillaring medium into the clay, (c) and heating the precursor at a temperature sufficient to decompose the added compound The pillared interlayered kandite compns. are suitable for use as catalysts, catalyst supports, sorbents, ion exchangers, extenders, fillers, and ceramic precursors.

IT 12026-57-2DP, inclusion compds. with halloysite RL: PREP (Preparation)

(manufacture of, for catalysts and sorbents and ceramic precursors)

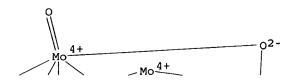
RN 12026-57-2 HCAPLUS

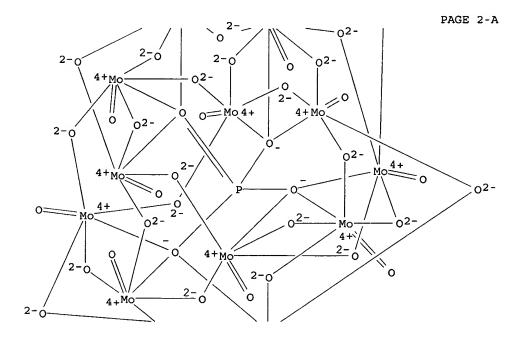
CN Molybdate(3-), tetracosa- μ -oxododecaoxo[μ 12-[phosphato(3-)- κ 0: κ 0: κ 0: κ 0': κ 0': κ 0'

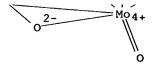
':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec

a-, trihydrogen (9CI) (CA INDEX NAME)

PAGE 1-A







●3 H+

IC ICM B01J029-02
CC 57-5 (Ceramics)
Section cross-reference(s): 67
IT Catalysts and Catalysis
Ceramic materials and wares
Cracking catalysts
Filling materials

Ion exchangers

Sorbents

(kandite for, pillared, intercalated, manufacture of)

1327-41-9DP, inclusion compds. with halloysite
12026-57-2DP, inclusion compds. with halloysite
12198-10-6DP, Garnierite, inclusion compds. with zirconium
oxychloride 12769-92-5DP, Zirconium chloride oxide, inclusion

compds. with halloysite RL: PREP (Preparation)

(manufacture of, for catalysts and sorbents and ceramic precursors)

L114 ANSWER 53 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1992:584843 HCAPLUS

DOCUMENT NUMBER:

117:184843

TITLE:

Polyoxotungstate compounds with antiviral

activity, especially against human

immunodeficiency resin (HIV)

INVENTOR(S):

Savage, Paul David; Theobald, Brian Ronald

Charles

PATENT ASSIGNEE(S):

Johnson Matthey PLC, UK

SOURCE: Eur. Pat. Appl., 17 pp.

CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

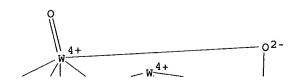
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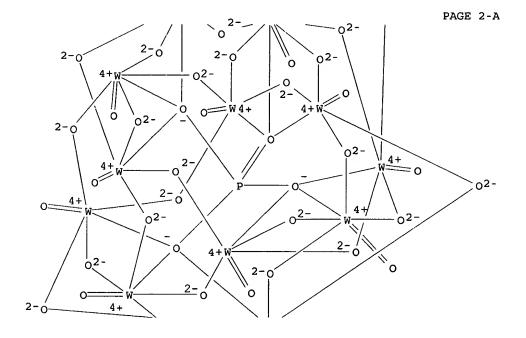
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EP 442663	A1	19910821	EP 1991-300997	
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				0206
	DE, DK	, ES, FR, G	B, GR, IT, LI, LU, NL,	SE
CA 2035994	AA	19910816	CA 1991-2035994	
				1991
				0208
AU 9170944	A1	19910822	AU 1991-70944	
				1991
				0208
ZA 9101065	Α	19911127	ZA 1991-1065	
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				0213
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				0215

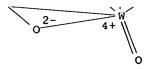
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PRIORITY APPLN. INFO.:
                                              GB 1990-3430
                                                                       1990
                                                                       0215
OTHER SOURCE(S):
                          MARPAT 117:184843
     Polyoxotungstate ions, having addnl. atoms selected from P,
     transition metals, and group 13 metals, and which have a metal
     cluster structure which is not the Keggin structure, form
     pharmaceutical compns. with antiviral, especially anti-HIV, activity. Examples of compds. of the invention include
     [P2HcW12Mx'Oy]P- (M' = transition metal; x = 0-6; y = 48-62; c =
     0-2; p = integer dependent on oxidation state of M'). Preparation of a
     large number of compds. of the invention is described, and activity
     of the compds. of the invention against HIV-1 and -2 is included.
ΙT
     85585-40-6
     RL: BIOL (Biological study)
        (antiviral)
DN
     85585-40-6 HCAPLUS
CN
     Vanadate(9-), [heptacosa-\u03c4-oxopentadecaoxo[\u03c49-[phosphato(3-)-
     κ0:κ0:κ0:κ0':κ0':κ0'':κ0
     '':κ0''':κ0''']]pentadecatungstate]nona-μ-
     oxotrioxo [μ9 - [phosphato (3-) -κ0:κ0:κ0:κ0]
     ':κ0':κ0'':κ0'':κ0''':κ0''']]tri-
     (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     101144-77-8
     RL: BIOL (Biological study)
        (antiviral activity against human immunodeficiency virus-1 and
        -2 of)
RN
     101144-77-8 HCAPLUS
     Vanadate (9-), [heptacosa-\mu-oxopentadecaoxo[\mu9-[phosphato(3-)-
     κ0:κ0:κ0:κ0':κ0':κ0'',κ0
     '':κ0''':κ0''']]pentadecatungstate]nona-μ-
     oxotrioxo [μ9-[phosphato(3-)-κ0:κ0:κ0:κ0
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*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
    1343-93-7 12027-38-2
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        (reaction of, in antiviral polyoxotungstate compound preparation)
     1343-93-7 HCAPLUS
CN
     Tungstate (3-), tetracosa-\mu-oxododecaoxo [\mu12-[phosphato(3-)-
     KO:KO:KO:KO':KO':KO':KO'
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a-, trihydrogen (9CI) (CA INDEX NAME)

PAGE 1-A



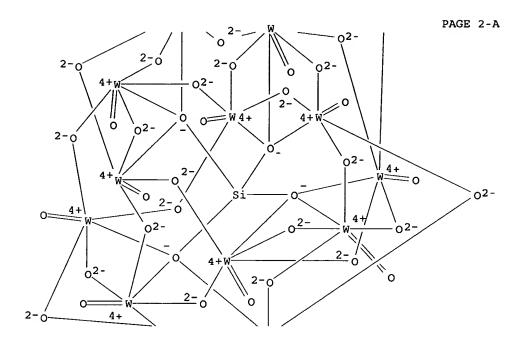


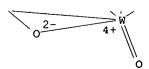


●3 H+

PAGE 1-A







●4 H+

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IC
     ICM A61K033-24
     ICS A61K033-42
CC
    1-5 (Pharmacology)
    Section cross-reference(s): 78
     85585-40-6
IT
                141216-02-6
    RL: BIOL (Biological study)
        (antiviral)
IT
    63950-53-8
                                                        79104-96-4
                 63950-56-1 65046-52-8
                                         79104-95-3
                 99397-47-4 101144-77-8 101347-05-1
     89173-95-5
    110294-54-7
                 110717-64-1 111933-31-4 139320-10-8, Cerium
    potassium tungsten oxide (CeK6W10O35)
                                            139320-11-9, Gadolinium
    potassium tungsten oxide (GdK7W10035)
                                            139320-12-0, Erbium
    potassium tungsten oxide (ErK7W10035)
                                            139381-06-9
                                                         139919-86-1.
    Praseodymium sodium tungsten oxide (PrNa7W10O35)
    RL: BIOL (Biological study)
        (antiviral activity against human immunodeficiency virus-1 and
       -2 of)
IT
    645-35-2 1343-93-7
                        7550-45-0, Titanium tetrachloride,
    reactions 7646-85-7, Zinc chloride (ZnCl2), reactions
    10098-89-2
                10141-05-6
                             10199-34-5
                                          11121-26-9, Silicotungstate
                              13568-40-6
                                           15595-35-4,
    12027-38-2
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    Arginine hydrochloride
                             52241-27-7
                                          53585-56-1 84750-84-5
    139083-47-9
    RL: RCT (Reactant); RACT (Reactant or reagent)
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APPLICATION NO.

DATE

(reaction of, in antiviral polyoxotungstate compound preparation)

L114 ANSWER 54 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

1992:536759 HCAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 117:136759

TITLE: Process using sorbents for removal of sulfur oxides from flue gas and other gas streams

Pinnavaia, Thomas J.; Amarasekera, Jayantha; INVENTOR(S):

DATE

Polansky, Christine A. Michigan State University, USA PATENT ASSIGNEE(S):

KIND

SOURCE: PCT Int. Appl., 38 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE:

English FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.

							_	DALL		TI I	DICHTION NO.			DAIL
							-							
	WO	91186	67			A1		199112	212	WO	1991-US3642			
														1991
														0528
		W:		•		D.	D.//	F0 F		an ar		0.77		
	110	KW: 51146		BE,	CH,						R, IT, LU, NL 1990-535147	, SE		
	03	31140	91			Α.		199203	17	US	1330-333147			1990
														0608
	CA	20647	15			AA		199112	209	CA	1991-2064715			
														1991
														0528
	ΕP	48667	6			A1		199205	27	EP	1991-912279			
														1991
	FD	48667	6			R 1		199610	100					0528
	D.									GB. GF	R, IT, LI, LU	. NI.	SF	:
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		07022						199503						
	AT	14382	7			E		199610	15	AT	1991-912279			1001
														1991 0528
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		10072				••		1,,,110	.23	C11	1331 103037			1991
														0608
PRIOR	(TIS	APPL	N.]	NFO.	. :					US	1990-535147	1	Ą	
														1990
														0608
										110	1990-466984	,	A 2	
										US	1990-400904			1990
														0118
										WO	1991-US3642	V	V	
														1991
														0528

The gases are contacted with a heated sorbent composition consisting of the layered double hydroxide structure [M1-x2+M'x3+(OH)2] (An-)x/n.yH2O where M, M' represent metal cations which form oxides and are capable of reacting with SO2 to form sulfites and with SO3 to form sulfates. A Is an interlayer oxo-anion that comprises ≥1 metal to provide oxidation of SO2 to SO3 (e.g., CrO42-, FeO42-, HVO42-, MoO42-, V100286-, Mo70246-, W70246-), and x = 0.8-0.12. Examples of the sorbents include

[Zn2Al(OH)6]NO3.zH2O and [Mg6Al2(OH)16]OH.xH2O, which are intercalated with anions such as V100286- and Keggin structures such as SiV3W9O407-. The sorbent can be added to the combustion chamber.

IT 143381-69-5 143381-71-9 143407-85-6 143407-87-8 143407-88-9 143414-73-7

RL: OCCU (Occurrence)

(sorbent, heated, for desulfurization of flue gases) 143381-69-5 HCAPLUS

RN

Tungstate (W12(OH)20386-), zinc (OC-6-11)-hexahydroxyaluminate(3-)
(1:12:6) (9CI) (CA INDEX NAME) CN

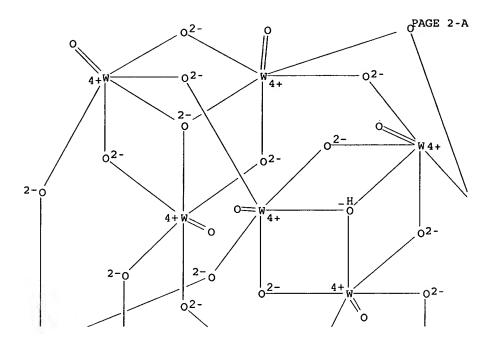
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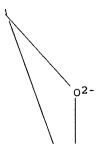
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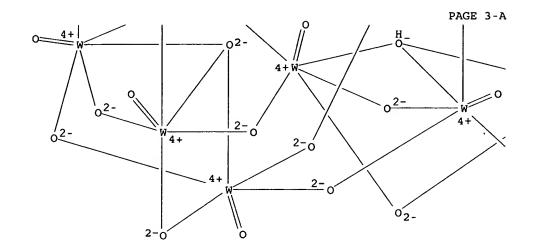
CRN 12207-61-3 CMF H2 O40 W12 CCI CCS

PAGE 1-A

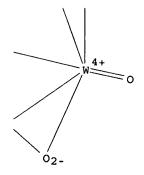


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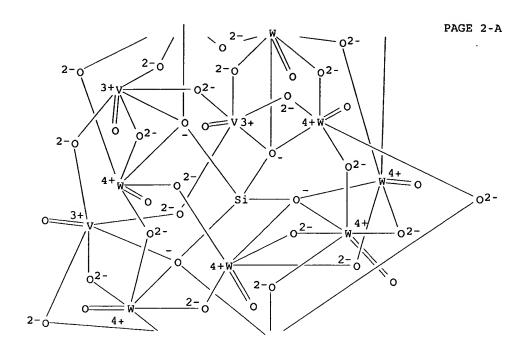


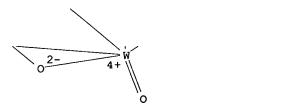
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CM 1

CRN 92816-60-9 CMF 040 Si V3 W9 CCI CCS





CM 2

CRN 18893-33-9 CMF Al H6 O6 CCI CCS

RN

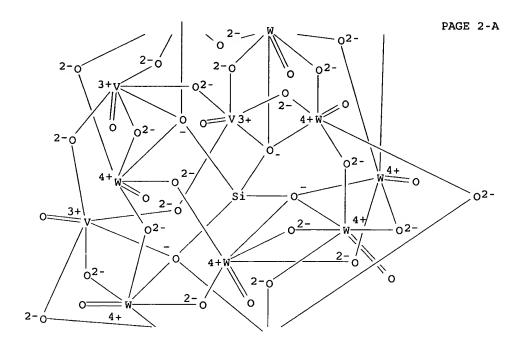
143407-85-6 HCAPLUS Vanadate(7-), [μ12-[orthosilicato(4-)-0:0:0:0':0':0'':0'':0''':0''':0''']]nona-μ-CN oxotrioxo(pentadeca-μ-oxononaoxononatungstate)tri-, magnesium (OC-6-11)-hexahydroxyaluminate(3-) hydroxide (1:21:7:14) (9CI) (CA INDEX NAME)

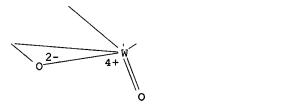
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CRN 92816-60-9 CMF 040 Si V3 W9

CCI CCS

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





PAGE 3-A

CM 2

CRN 18893-33-9 CMF Al H6 O6 CCI CCS

RN 143407-87-8 HCAPLUS

CN Tungstate (W12(OH)20386-), magnesium (OC-6-11)hexahydroxyaluminate(3-) hydroxide (1:18:6:12) (9CI) (CA INDEX
NAME)

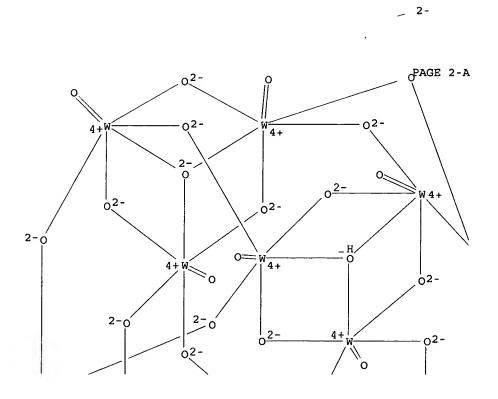
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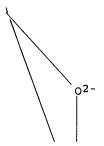
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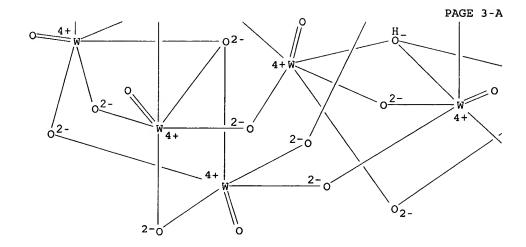
CRN 12207-61-3 CMF H2 O40 W12 CCI CCS

PAGE 1-A

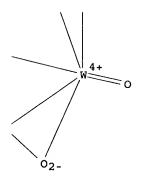


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PAGE 3-B

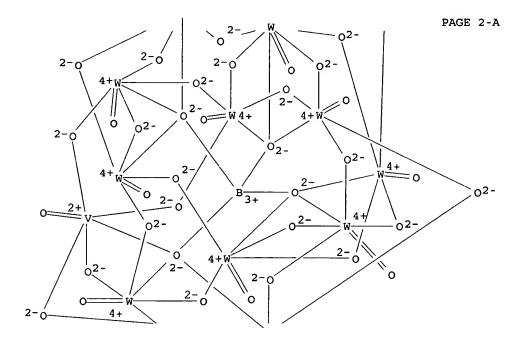


RN 143407-88-9 HCAPLUS
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(OC-6-11)-hexahydroxyaluminate(3-) hydroxide (1:21:7:14) (9CI)
(CA INDEX NAME)

CM 1

CRN 53260-17-6
CMF B 040 V W11
CCI CCS

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





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CRN 18893-33-9 CMF Al H6 O6 CCI CCS

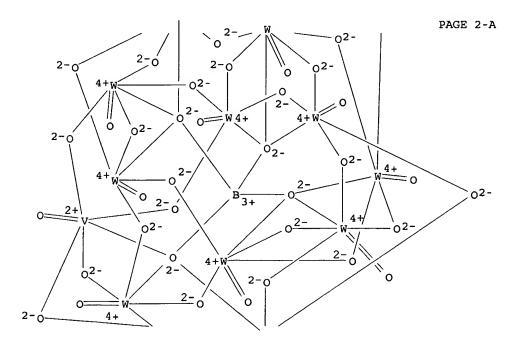
RN 143414-73-7 HCAPLUS

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(OC-6-11)-hexahydroxyaluminate(3-) (1:14:7) (9CI) (CA INDEX NAME)

CM 1

CRN 53260-17-6 CMF B O40 V W11

CCI CCS



02- W

CM

PAGE 3-A

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CRN 18893-33-9
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         CCS
     CCI
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      OH-
IC
     ICM B01J008-00
     ICS C01B017-00; C09K003-00
59-4 (Air Pollution and Industrial Hygiene)
CC
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TΤ
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     143480-83-5
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         (sorbent, heated, for desulfurization of flue gases)
L114 ANSWER 55 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                          1991:688014 HCAPLUS
DOCUMENT NUMBER:
                          115:288014
TITLE:
                          Negative selectivity in reverse-osmotic
                          separation of heteropoly anions and nitrate
                          ions in an aqueous solutions
AUTHOR(S):
                          Semin, G. L.; Vetchinova, Yu. S.; Il'nich, O.
                          M.; Fedotov, M. A.; Kuznetsova, L. I.;
                          Novopashina, V. M.
CORPORATE SOURCE:
                          Inst. Catal., Novosibirsk, USSR
SOURCE:
                          Khimiya i Tekhnologiya Vody (1991), 13(8),
                          685-8
                          CODEN: KTVODL; ISSN: 0204-3556
DOCUMENT TYPE:
                          Journal
LANGUAGE:
                          Russian
     The characteristics of composite polyamide membranes
     OPM-K, OPAM-K and OPMN-K (Scientific Industrial
     Amalgamation Polymersintez», Vladimir) relative to retention to the Na salt of the heteropolyacid (HPA) catalyst
     Na4PW11Fe3+(H2O)O39 and NaNO were determined as functions of pH,
     pressure, and temperature Expts. on separation of a mixture of these
     salts were performed. The selectivity of heteropolyanion
     retention greatly exceeds the selectivity of nitrate retention;
```

this permits efficient separation of HPA from NO3-. The selectivity of nitrate retention from a mixture with HPA depends on the HPA concn; with increasing HPA concentration, the selectivity decreases and attains neg. values.

IT 137827-83-9

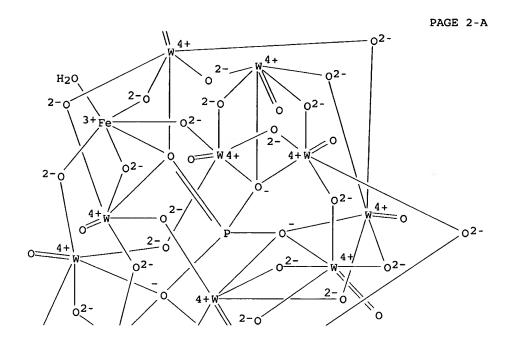
RL: PRP (Properties)
(reverse osmosis separation of, on polyamide membranes)

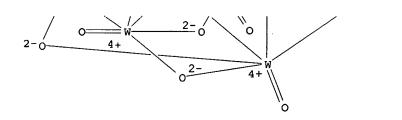
RN 137827-83-9 HCAPLUS

Tungstate(4-), (aquaferrate)tetracosa-μ-oxoundecaoxo(μ12[phosphato(3-)-κΟ:κΟ:κΟ:κΟ':.kap
pa.O':κΟ'':κΟ'':κΟ'':.ka
ppa.O''']]undeca-, tetrasodium (9CI) (CA INDEX NAME)

PAGE 1-A

Les Henderson Page 416 571-272-2538





•4 Na+

CC 66-4 (Surface Chemistry and Colloids)

Section cross-reference(s): 38, 48, 59, 60

IT 137827-83-9

RL: PRP (Properties)

(reverse osmosis separation of, on polyamide membranes)

L114 ANSWER 56 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1991:670659 HCAPLUS

DOCUMENT NUMBER:

115:270659

TITLE:

Pharmaceutical composition

containing polyoxymetallate compounds for AIDS

treatment

INVENTOR(S):

Murrer, Barry Anthony; Theobald, Brian Ronald

Charles; Savage, Paul David Johnson Matthey PLC, UK

PATENT ASSIGNEE(S): SOURCE:

Eur. Pat. Appl., 7 pp.

CODEN: EPXXDW

DOCUMENT TYPE: LANGUAGE:

Patent English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT N	0.	KIND	DATE	APPLICATIO	ON NO.	DATE
EP 39036	5		19901003	EP 1990-30	02721	1000
						1990 0314
R: . CA 20121				GB, GR, IT, I CA 1990-20		SE
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HU 53655		A2	19901128	HU 1990-19	593	1990
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HU 20733			19930329			
US 50931	34	А	19920303	US 1990-49	93342	1990
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ZA 90019	93	A	19910130	ZA 1990-19	993	
						1990 0315
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AO	10	AI	15500520	AU 1330 3.	1410	1990
	_					0316
AU 62277 JP 03047		B2 A2		JP 1990-64	1457	
UP 03047	130	AZ	19910220	OF 1990-64	1457	1990
						0316
PRIORITY APPL	N. INFO.:			GB 1989-61	189	A 1989
						0317

AB Polyoxymetallate compds. Ax[MDLW11039].yH20 (I) and Ax[D'(MW11039)2].yH2O (A = cation; x = integer varying with M and oxidation state of D or D'; M = B, Si, P; D = metal; D' = lanthanide in oxidation state 3 or 4; L = neutral or anionic ligand; y = integer; provided that when D = Co, V, or Al, L ≠ H2O) are in pharmaceutical compns. for the treatment of human immunodeficiency virus (HIV)-infected patients or prophylactic treatment of patients at risk from HIV infection.

K15[Er(BW11039)2].yH2O was prepared by reacting Na2WO4.2H2O and boric acid, and treating the [BW11039]9- species produced with Er(NO3)3.5H2O. Compds. I exhibited selective activity against HIV-1 and HIV-2 in infected cells and their toxicity was much less than that of HPH-23 (NaSb9W21086). For example, K13[Ce(SiW11039)2].26H2O had a CD50 (50% cytotoxic dose) of >1000.0 μg/mL, an antiviral ED50 value of 0.39 μg/mL, and a selectivity index (SI = CD50/ED50) of >2561 against HIV-1 in infected MT-4 cells.

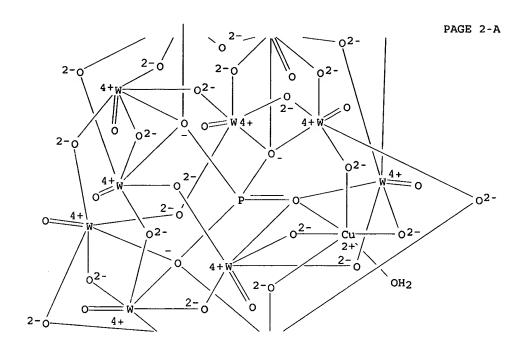
IT 66257-58-7D, salt hydrates 66304-44-7D, salt hydrates 66304-53-8D, salt hydrates 81552-96-7 81553-01-7D, salt hydrates 81553-37-9D, salt hydrates 135143-89-4D, salt hydrates 135211-05-1D, salt hydrates 136292-63-2D, salt hydrates 136314-55-1D, salt hydrates RL: BIOL (Biological study)

(AIDS treatment with)

RN 66257-58-7 HCAPLUS

CN Tungstate(5-), (aquacuprate)tetracosa-μ-oxoundecaoxo[μ12[phosphato(3-)-κ0:κ0:κ0':κ0':.kap
pa.0':κ0'':κ0'':κ0'':κ0'':.ka
ppa.0'']]undeca-, pentahydrogen (9CI) (CA INDEX NAME)

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





●5 H+

RN 66304-44-7 HCAPLUS

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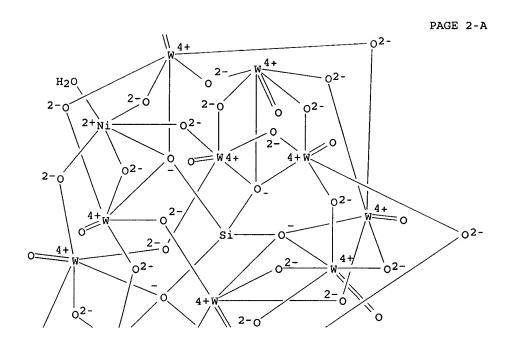
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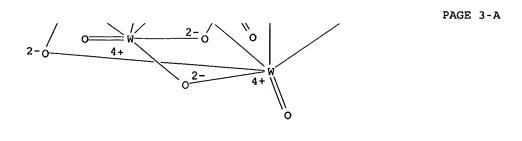
RN 66304-53-8 HCAPLUS

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PAGE 1-A

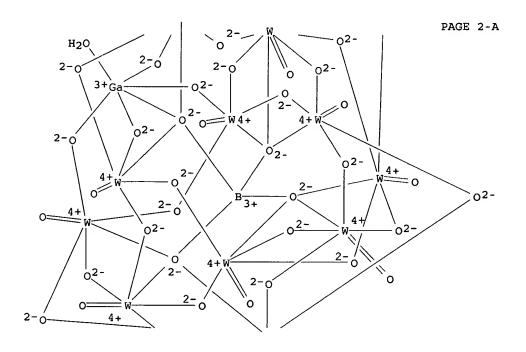






●6 H+

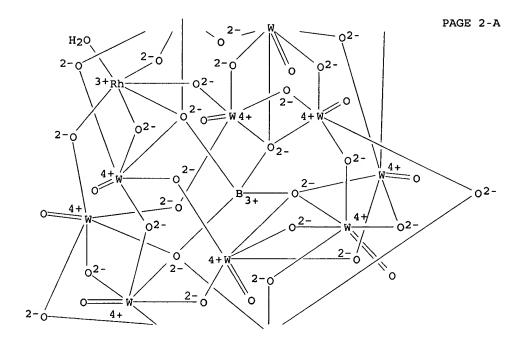
RN 81552-96-7 HCAPLUS
CN Tungstate(6-), (aquagallate)tetracosa-µ-oxoundecaoxo[µ12[tetrahydroxyborato(5-)-0:0:0':0':0'':0'':0'':0''':0''']
]undeca-, hexahydrogen (9CI) (CA INDEX NAME)

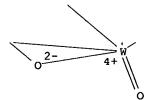




●6 H⁴

RN 81553-01-7 HCAPLUS
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]undeca-, hexahydrogen (9CI) (CA INDEX NAME)





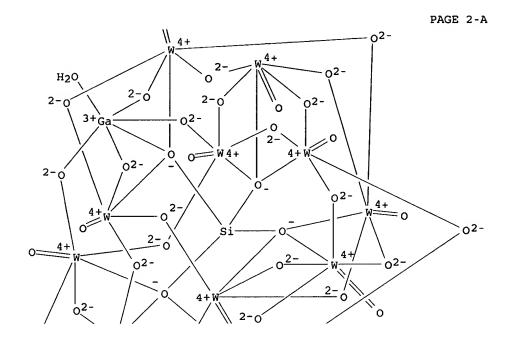
●6 H+

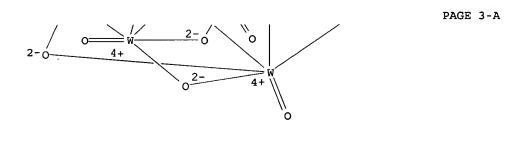
RN 81553-37-9 HCAPLUS

CN Tungstate(5-), (aquagallate) [μ12-[orthosilicato(4-)Ο:Ο:Ο:Ο':Ο':Ο'':Ο'':Ο''':Ο''']]tetracosa-μoxoundecaoxoundeca-, pentahydrogen (9CI) (CA INDEX NAME)

PAGE 1-A

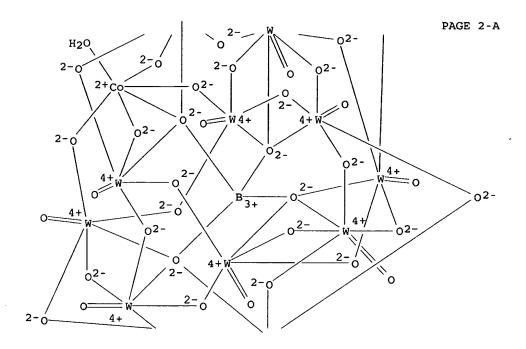
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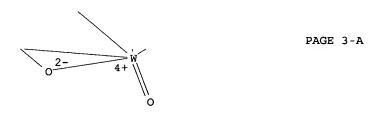




●5 H+

RN 135143-89-4 HCAPLUS
CN Tungstate(7-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-0:0:0':0':0'':0'':0'':0''':0''']
] undeca-, heptahydrogen (9CI) (CA INDEX NAME)



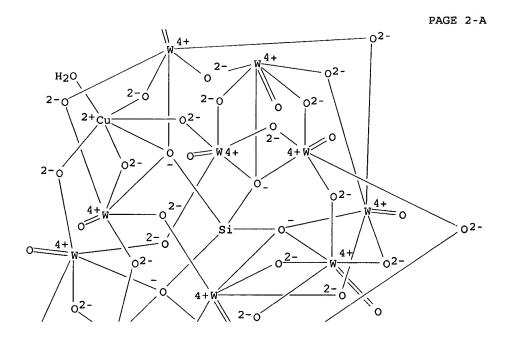


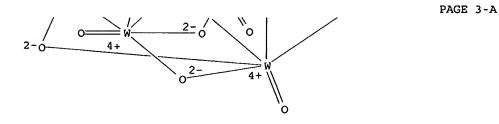
●7 H+

RN 135211-05-1 HCAPLUS CN Tungstate(6-), (aquacuprate)[μ 12-[orthosilicato(4-)-0:0:0:0':0':0'':0'':0'':0''':0''']]tetracosa- μ -oxoundecaoxoundeca-, hexahydrogen (9CI) (CA INDEX NAME)

PAGE 1-A

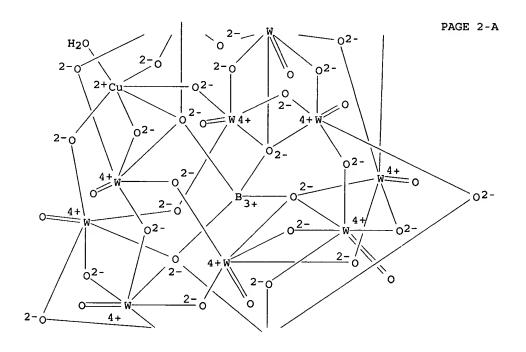






●6 H+

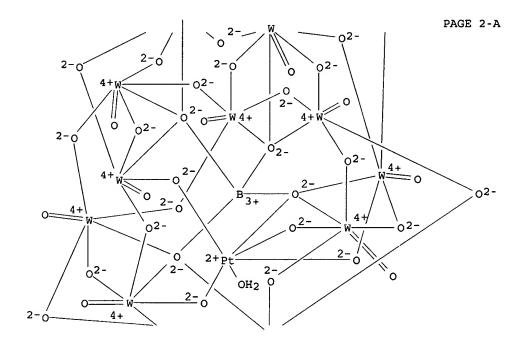
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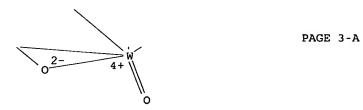




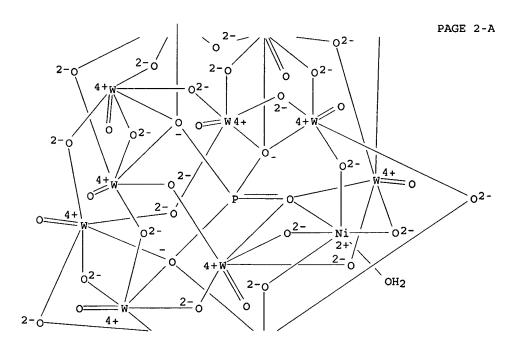
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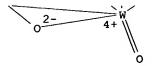
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●7 H+

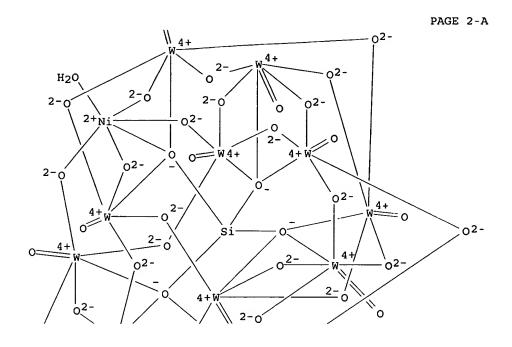


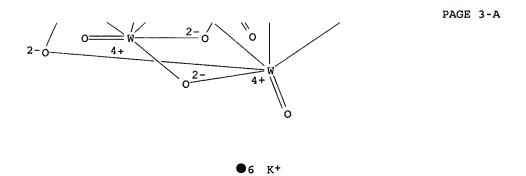


●5 K+

PAGE 1-A

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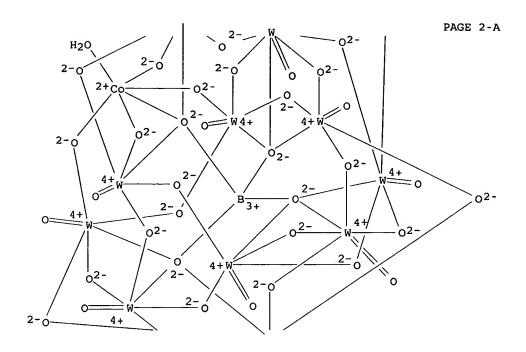


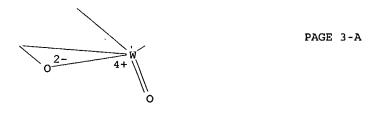


RN 39291-87-7 HCAPLUS
CN Tungstate(7-), (aquacobaltate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ::.kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':kappa
.O''':κΟ''']]undeca-, heptapotassium (9CI) (CA INDEX NAME)

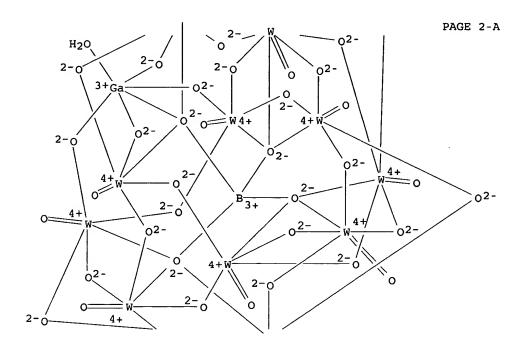
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT

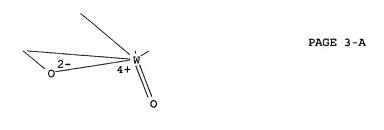
Les Henderson





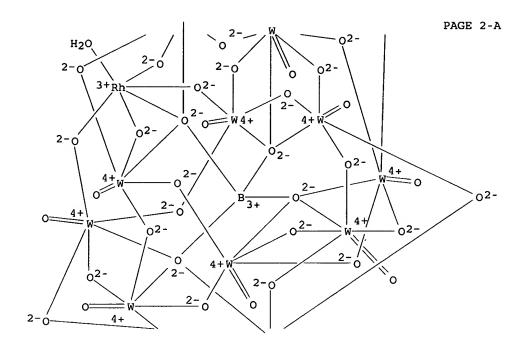
●7 K+

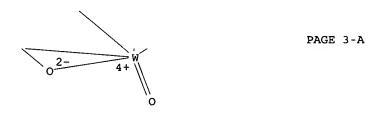




●6 K+

RN 81553-02-8 HCAPLUS
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]undeca-, hexapotassium (9CI) (CA INDEX NAME)

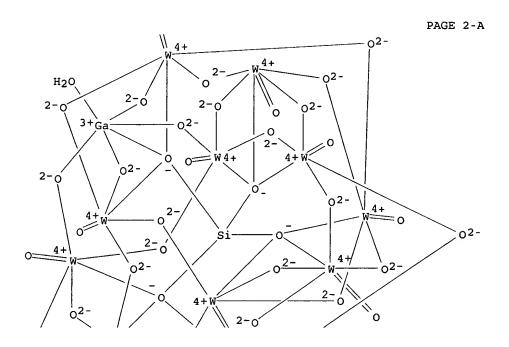


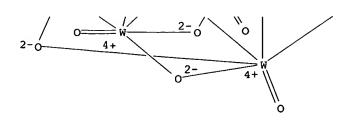


●6 K+

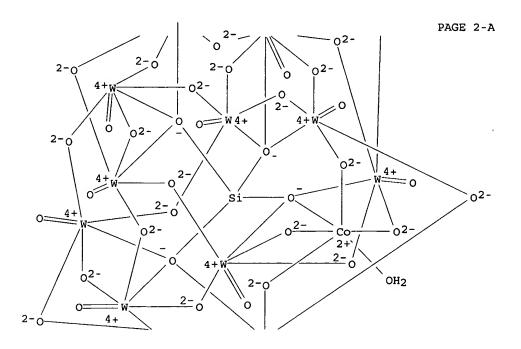
RN 81553-38-0 HCAPLUS
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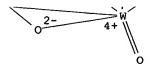






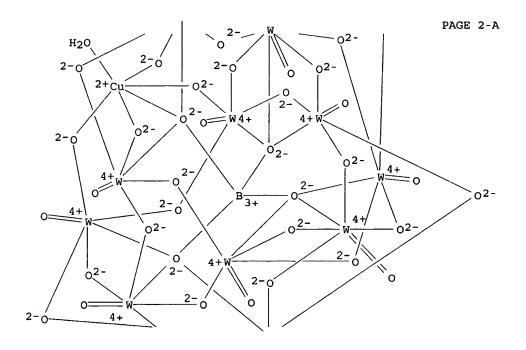
●5 K+

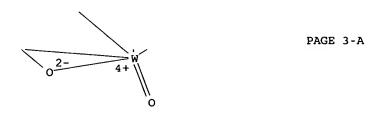




●6 K+

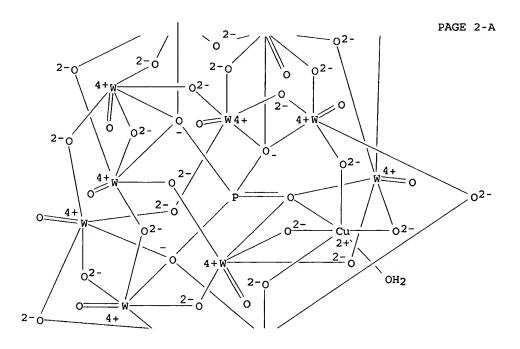
RN 135143-90-7 HCAPLUS
CN Tungstate(7-), (aquacuprate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ:κΟ::.kapp
a.O':κΟ':κΟ'':κΟ'':κΟ'':.kappa
.O''':κΟ''']]undeca-, heptapotassium (9CI) (CA INDEX NAME)

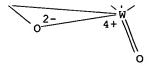




●7 K+

RN 135244-69-8 HCAPLUS
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ppa.O''']]undeca-, pentapotassium (9CI) (CA INDEX NAME)

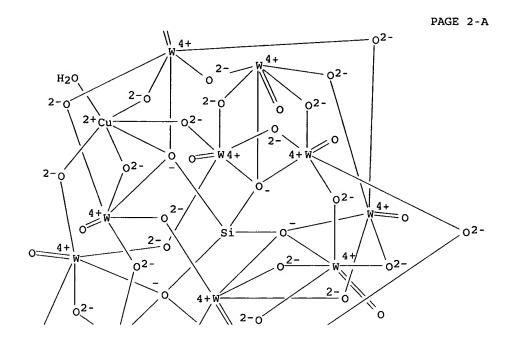


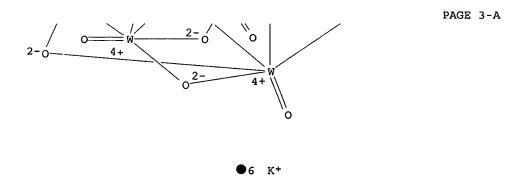


●5 K⁺

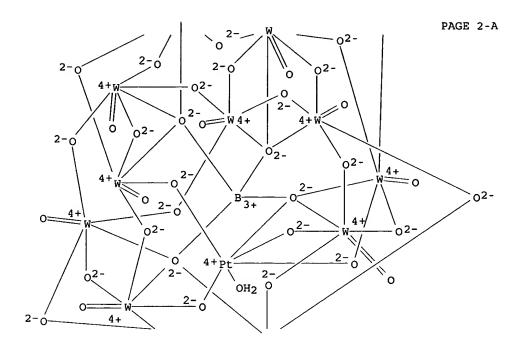
PAGE 1-A

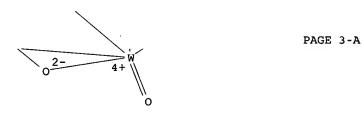
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RN 137679-48-2 HCAPLUS
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]undeca-, pentapotassium (9CI) (CA INDEX NAME)





●5 K+

```
ICM A61K033-24
IC
     ICS A61K033-42
CC
     1-5 (Pharmacology)
     Section cross-reference(s): 63, 78
IT
     66257-58-7D, salt hydrates 66304-44-7D, salt
     hydrates 66304-53-8D, salt hydrates 81552-96-7
     81553-01-7D, salt hydrates 81553-37-9D, salt
     hydrates 108174-18-1D, salt hydrates 135143-89-4D,
     salt hydrates 135211-05-1D, salt hydrates
     135244-67-6D, salt hydrates 135500-08-2D, salt hydrates 135523-13-6D, salt hydrates 136292-63-2D, salt hydrates
     136314-55-1D, salt hydrates
     RL: BIOL (Biological study)
        (AIDS treatment with)
     37194-75-5 37194-76-6 39291-87-7
IT
     81552-97-8 81553-02-8 81553-38-0
     105785-76-0 106096-53-1 135143-90-7
     135244-68-7 135244-69-8 135266-66-9
     135596-17-7 135635-38-0 137679-48-2
     RL: BIOL (Biological study)
         (human immunodeficiency virus infection treatment with)
```

L114 ANSWER 57 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

115:261849

ACCESSION NUMBER: DOCUMENT NUMBER:

1991:661849 HCAPLUS

TITLE:

Ceramic compositions for insulating

parting layers in two-stage precision casting

and their manufacture

INVENTOR(S):

Wall, Giselher

PATENT ASSIGNEE(S):

Germany

SOURCE:

Ger. Offen., 9 pp. Addn. to Ger. Offen.

3,903,427. CODEN: GWXXBX

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 4001057	A1	19910718	DE 1990-4001057	
				1990
				0116
DE 4001057	C2	19960711		
DE 3903427	A1	19900809	DE 1989-3903427	
				1989
DT 2002405		10001105		0206
DE 3903427	C2	19921105	DD 1000 337661	
DD 293972	A 5	19910919	DD 1990-337661	1000
				1990
PRIORITY APPLN. INFO.:			DE 1989-3903427	0207
PRIORITY APPLIN. INFO.:			DE 1989-3903427	1989
				0206
				0200
			DE 1990-4001057 A	
				1990
				0116

AB The parting layer, mech. applied to the primary part and then sintered, comprise (a) ≥1 alkaline earth oxides, a rare earth oxide, Al203 or a Ti-group oxide, or (b) ≥1 double oxides, or a mixture of a metal oxide and a nonmetal oxide, e.g., B203, P205, or SiO2, and corresponding compds. in which the O is replaced by a halogen, especially F. The layers are prepared from ceramic compns. containing high-melting particles consist of an oxide as above or SiC, SiO2 fume, or a nitride and low-melting particles of stoichiometrically defined double or mixed oxides and correspond to an anhydrous salt of silicic, phosphoric, boric, or metal acid, including halogen-free heteropolyacids and Keggin acids, in which O may be replaced by halogen, preferably F, and the cation is an alkali metal ion, preferably K. The 2-stage precision casting method is especially suitable for the manufacture of dental material anchors, e.g., double crowns, and jewelry. Parting layers, prepared from TiO2 and red Fe oxide in 50% aqueous H3PO4 containing .apprx.5% glycol were resistant against Au alloys.

IT 113857-50-4, Tungstosilicic acid

RL: USES (Uses)

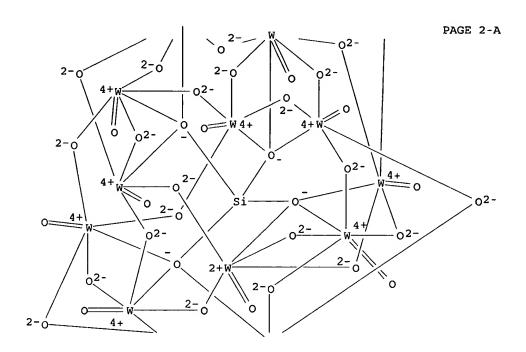
(in ceramic parting layer manufacture)

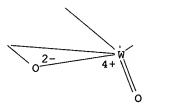
113857-50-4 HCAPLUS RN

Tungstate(6-), $[\mu 12$ -[orthosilicato(4-)-CN

0:0:0:0':0':0':0'':0'':0'':0''':0''']]tetracosa-μ-oxododecaoxododeca-, hexahydrogen (9CI) (CA INDEX NAME)

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





PAGE 3-A

●6 H+

IC ICM A61K006-06
ICS A61C013-20; A61C013-00; B22C003-00
CC 57-2 (Ceramics)
Section cross-reference(s): 56, 63
IT Alkali metal oxides
Aluminates
Ferrates
Rare earth oxides
Stannates
Tungstates
Vanadates
Zirconates
RL: USES (Uses)

(ceramic parting layer compns. containing, for two-stage precision casting)

IT Ceramic materials and wares

(parting layers, in two-stage precision casting of dental materials and jewelry)

```
TΤ
     Jewelry and Jewels
        (precision casting of, two-stage, ceramic compns. for
        parting layer in secondary casting in)
     Dental materials and appliances
TΤ
        (alloys, crowns, precision casting of, two-stage, ceramic
        compns. for parting layer in secondary casting in)
IT
     Group IVB element chalcogenides
     RL: USES (Uses)
        (oxides, ceramic parting layer compns. containing, for
        two-stage precision casting)
TT
     Casting process
         (precision, two-stage, ceramic compns. for parting
        layer in secondary casting in, for dental materials and
     137442-83-2
                    137442-84-3 137442-85-4
TΤ
                                                137442-86-5
     RL: PEP (Physical, engineering or chemical process); PROC
     (Process)
        (casting of, precision, ceramic parting layer compns.
        for, for dental materials and jewelry)
IT
     1303-86-2, Boron oxide, uses and miscellaneous
                                                        1309-37-1, Iron
     oxide (Fe2O3), uses and miscellaneous 1314-56-3, Phosphorus
     pentoxide, uses and miscellaneous 7631-95-0, Sodium molybdate 10043-11-5, Boron nitride, uses and miscellaneous 12033-89-5,
     Silicon nitride, uses and miscellaneous 12705-37-2, Chromium
               13721-39-6, Sodium orthovanadate 13775-52-5, Potassium
     hexafluoroaluminate 16919-27-0 24304-00-5, Aluminum nitride
     24646-85-3, Vanadium nitride 25583-20-4, Titanium nitride
     RL: USES (Uses)
        (ceramic parting layer compns. containing, for two-stage
        precision casting)
     56-81-5, 1,2,3-Propanetriol, uses and miscellaneous
IT
               79-39-0, Methacrylic acid amide 107-21-1,
     Pinacol
     1,2-Ethanediol, uses and miscellaneous 7664-38-2, Phosphoric acid, uses and miscellaneous 9003-39-8, Polyvinylpyrrolidone
     11104-88-4, Molybdophosphoric acid
                                           42615-58-7, Tungstoboric acid
     113857-50-4, Tungstosilicic acid
     RL: USES (Uses)
        (in ceramic parting layer manufacture)
L114 ANSWER 58 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                          1987:412131 HCAPLUS
DOCUMENT NUMBER:
                          107:12131
TITLE:
                          Characteristics of catalysts for
                          low-temperature carbon monoxide oxidation
                          obtained by supporting palladium and
                          heteropolyacid on silica gel
AUTHOR(S):
                          Pavlova, S. N.; Kuznetsova, L. I.; Matveev, K.
                          I.; Sazonov, V. A.; Popovskii, V. V.;
Zhizhina, E. G.; Fenelonov, V. B.; Gavrilov,
                          V. Yu.
CORPORATE SOURCE:
                          Inst. Katal., Novosibirsk, USSR
SOURCE:
                          Kinetika i Kataliz (1987), 28(2), 373-9
                          CODEN: KNKTA4; ISSN: 0453-8811
DOCUMENT TYPE:
                          Journal
LANGUAGE:
                          Russian
     A catalyst consisting of Pd supported on silica gel saturated with the
     aqueous phase of heteropolyacid (composition H7PMo8V4O40.30H2O)
     was effective for the oxidation of CO in moist waste gases. The
     oxidation expts. were conducted at 15-30° and 35-100% relative
     humidity of the waste gases. The maximum CO oxidation (82%) was
     obtained when the catalyst contained 1.37% Pd and 7.8%
     heteropolyacid. The catalyst was a 2-phase system consisting of
     Pd on solid silica gel and heteropolyacid in aqueous phase held by the
     silica gel pores.
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Les Henderson Page 444 571-272-2538

TТ

104574-43-8

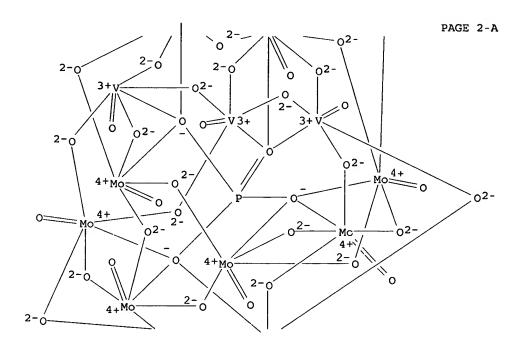
RL: OCCU (Occurrence)

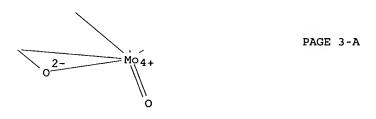
(in oxidation catalysts containing palladium, on silica gel, for carbon monoxide removal from moist waste gases)

RN 104574-43-8 HCAPLUS

CN

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT





●7 H+

●x H2O

CC 59-4 (Air Pollution and Industrial Hygiene)
 Section cross-reference(s): 67

IT 104574-43-8

RL: OCCU (Occurrence)

(in oxidation catalysts containing palladium, on silica gel, for carbon monoxide removal from moist waste gases)

L114 ANSWER 59 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 1986:11207 HCAPLUS DOCUMENT NUMBER: 104:11207

TITLE: Analytical use of solvent extraction with

acetonitrile/water/chloroform and 1-propanol/water/cyclohexane mixtures

AUTHOR(S):

Hori, T.; Fujinaga, T. Coll. Lib. Arts Sci., Kyoto Univ., Kyoto, 606, CORPORATE SOURCE:

Japan

SOURCE: Talanta (1985), 32(8B), 735-43

CODEN: TLNTA2; ISSN: 0039-9140

DOCUMENT TYPE: Journal LANGUAGE: English

To obtain an organic phase containing MeCN or 1-PrOH from aqueous MeCN or aqueous PrOH solution, the effect of CHCl3 and cyclohexane as auxiliary solvents was investigated. Use of such ternary system offers an alternative to the so-called salting-out method, in which inorg. salts or hydrophilic organic materials are used to sep. organic phases from otherwise miscible solvent mixts. It also seems preferable for solvent-extraction procedures, since only a small amount of auxiliary solvent is needed instead of usually large amount of salt (impurities in which may cause undue contamination); also, the volume and composition of the organic phase can be predicted from phase diagrams and the overall composition of the solvent mixture Volume-fraction diagrams are easy to use. Furthermore, equilibrium is attained in solvent mixts. more rapidly than in salting-out systems. The utility of the ternary solvent systems was demonstrated for extraction of intermediate molybdophosphate complexes which are formed in aqueous acetonitrile solns

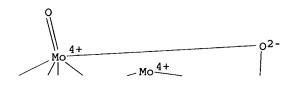
ΙT 12379-13-4

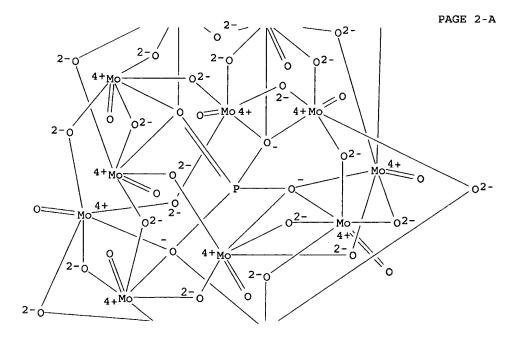
RL: PROC (Process)

(extraction of, by acetonitrile-chloroform-water mixture)

12379-13-4 HCAPLUS RN

Molybdate (3-), tetracosa- μ -oxododecaoxo [μ 12-[phosphato(3-)-CN INDEX NAME)





CC 68-2 (Phase Equilibriums, Chemical Equilibriums, and Solutions) 12379-13-4 14526-22-8 IT

RL: PROC (Process)

(extraction of, by acetonitrile-chloroform-water mixture)

L114 ANSWER 60 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1983:433979 HCAPLUS

DOCUMENT NUMBER:

99:33979

TITLE: Spot tests for cannabis materials

Ι

Tewari, S. N.; Sharma, J. D.

AUTHOR (S): CORPORATE SOURCE:

Forensic Sci. Lab., Lucknow, India Bulletin on Narcotics (1982), 34(3-4), 109-12 SOURCE:

CODEN: BNUNA5; ISSN: 0007-523X

DOCUMENT TYPE: Journal

LANGUAGE: English

GI

Eight spot tests were introduced for the identification of AΒ different cannabis drugs. They consisted in testing the suspecting materials with gel layer or filter paper strips impregnated with chromogenic substances [for example p-aminoantipyrine (I) [69267-59-0], tetrazolium blue [1871-22-3], etc.]. All the tests gave satisfactory results especially when gel layers were used as the reaction surfaces. The test which used cinnamaldehyde [104-55-2] as the chromogenic agent was the most sensitive, it detected 0.1, 0.5, and 2.0 $\mu g/mL$ of the cannabis species resin, ganja, and bhang, resp. IT

12026-57-2

RL: BIOL (Biological study)

(cannabis materials detection by, in spot test)

RN 12026-57-2 HCAPLUS

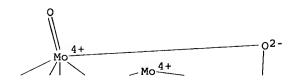
CN Molybdate(3-), tetracosa- μ -oxododecaoxo[μ 12-[phosphato(3-)-

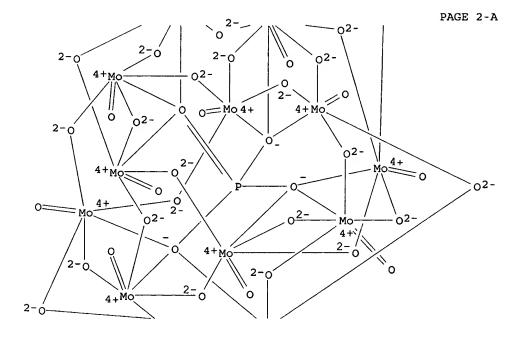
κ0:κ0:κ0:κ0':κ0':κ0':κ0'

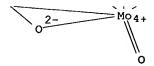
':κ0'':κ0'':κ0''':κ0''':κ0''']]dodec

a-, trihydrogen (9CI) (CA INDEX NAME)

PAGE 1-A







CC 4-2 (Toxicology)

103-84-4 104-55-2 108-24-7 123-30-8 366-18-7 1310-58-3, biological studies 1310-73-2, biological studies 1871-22-3 TΤ 7664-93-9, biological studies 12026-57-2 27766-45-6 69267-59-0

RL: BIOL (Biological study)

(cannabis materials detection by, in spot test)

L114 ANSWER 61 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

1983:217344 HCAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 98:217344

TITLE: Fluororesin-containing coating

composition

Yoshimura, Tatsushiro; Tominaga, Shigetake INVENTOR(S):

Daikin Kogyo Co., Ltd. , Japan Eur. Pat. Appl., 22 pp. PATENT ASSIGNEE(S):

SOURCE:

CODEN: EPXXDW

DOCUMENT TYPE:

Patent English

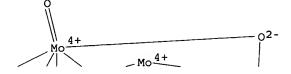
LANGUAGE:

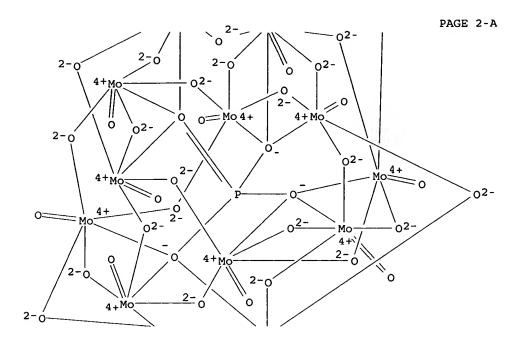
FAMILY ACC. NUM. COUNT:

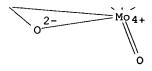
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 75908	A1	19830406	EP 1982-108901	
				1982
				0925
EP 75908	B1	19841227		
R: DE, FR, GB				
JP 58053960	A2	19830330	JP 1981-154180	
				1981
				0928
JP 61016293	B4	19860430		
US 4521596	A	19850604	US 1982-423189	
				1982
				0924
PRIORITY APPLN. INFO.:			JP 1981-154180 A	0,22,1
			1 1301 131100 11	1981
				0928
				UフZB

AB Fluorocarbon polymer coatings are adhered to metal surfaces using primer coatings containing fluororesin dispersion, H2MoO4 or its salt, NH4OH, H3PO4 or its salt, and colloidal SiO2, with the coating having good heat and corrosion resistance and no toxicity or pollutant release. Thus, 16 parts MoO3 was dissolved in 40 parts 28% aqueous NH4OH and diluted with 50 parts water. parts 85% H3PO4 was added followed by 100 parts colloidal SiO2. Then, the composition was mixed with polytetrafluoroethylene [9002-84-0] (55% dispersion) in ratio 20:10. The resulting primer was sprayed on an iron plate, dried 10 min at 100° and baked 20 min at 380°. A topcoat of hexafluoropropylene-tetrafluoroethylene copolymer [25067-11-2]







●3 NH4+

C09D003-78; C09D005-08 IC

42-10 (Coatings, Inks, and Related Products) CC

Section cross-reference(s): 55, 56

IT Coating materials

> (perfluorocarbon polymer, adhesion of, to metal surfaces, compns. for)

IT 9002-84-0 25067-11-2

RL: TEM (Technical or engineered material use); USES (Uses) (coatings, for metal surfaces, adhesion of, compns.

ΙT 1313-27-5, uses and miscellaneous 1336-21-6 7631-86-9, uses and miscellaneous 7664-38-2, uses and miscellaneous 7782-91-4 12026-66-3 12027-67-7 34085-20-6 61583-60-6

RL: USES (Uses)

(primer coating compns. containing, for adhering perfluorocarbon polymer coatings to metals)

L114 ANSWER 62 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1975:526488 HCAPLUS

DOCUMENT NUMBER: 83:126488

TITLE: Complex tungsten compounds and their

pharmaceutical use

INVENTOR(S): Chermann, Jean C.; Jasmin, Claude; Mathe,

Georges; Raynaud, Marcel

PATENT ASSIGNEE(S):

Agence Nationale de Valorisation de la

Recherche

SOURCE:

Fr. Demande, 47 pp. Division of Fr. Demande

2,117,803 (See Ger. 2,162,373, CA 77:

143794n). CODEN: FRXXBL

DOCUMENT TYPE: LANGUAGE:

Patent French

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
FR 2161837	A2	19730713	FR 1971-43444	1971
FR 2161837 PRIORITY APPLN. INFO.:	B2	19750207	FR 1971-43444 A	1203 1971
				1203

AB A series of 20 complex heteropolyanionic compds. (I), e.g., P2W18O62(NH4)6 [37300-90-6], SiW11O39CrIIIH5 [56367-35-2], PV2Mo10040H5 [12293-21-9] and AsMo12040H7 [56449-73-1], are useful for treatment of diverse viral infections. I consist of a plurality of anions derived from weak acids such as tungstic, molybdic and vanadic acids, grouped around a central atom such as B, Si, Ge, P, or As through coordination bonds with W, Mo, or V, in a ratio of W or Mo/central atom ranging mainly 8.5-12. One or more of the W or Mo atoms may also be substituted by V, Cr, Fe, Co, Ni, or Cu. Aqueous solns. of a precursor or Na salt of the central atom and of a precursor or Na salt of the weak acid, in definite molar concns. as desired, are mixed in the presence of a concentrated acid such as 5N H2SO4; the resulting solution is boiled under reflux for 5 hr and allowed to cool, and the precipitate formed is filtered to give the desired I in acid form. Ammonium, alkali metal, and alkaline earth salts are prepared by neutralization of the acid form with appropriate hydroxides. The preparation of I in which ≥ 1 W atoms are substituted by a metal is described. To a boiling solution of silico-12 tungstic acid in H2O-AcOH, adjusted to pH 6 with KHCO3, is added an aqueous solution of Co(OAc)2 and a solution of KOAc in H2O-AcOH; the reaction mixture is filtered and kept overnight at 5°, and the deposited crystals are filtered and recrystd. from warm water to give I in which W is substituted by Co2+. Preclin. pharmacol. data are included, especially for SiW12040Na4 [12027-47-3]. Some of the compds. inhibited several viruses in vitro and in vivo.

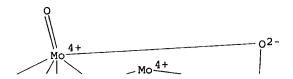
12026-57-2 12027-38-2 12027-47-3 TT 12293-21-9 12297-12-0 37300-77-9 37306-30-2

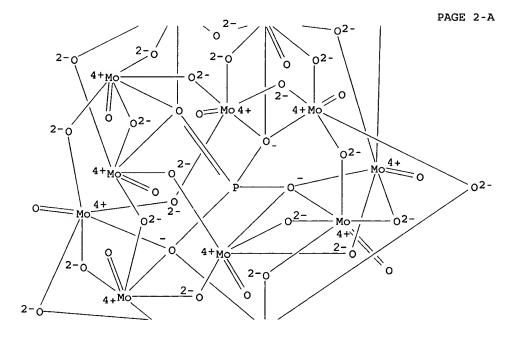
> RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study) (virucide)

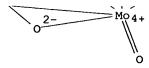
RN 12026-57-2 HCAPLUS

Molybdate(3-), tetracosa- μ -oxododecaoxo[μ 12-[phosphato(3-)κ0:κ0:κ0:κ0':κ0':κ0':κ0' ':κ0'':κ0'':κ0''':κ0''']]dodec a-, trihydrogen (9CI) (CA INDEX NAME)

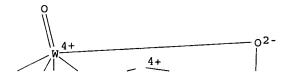
PAGE 1-A

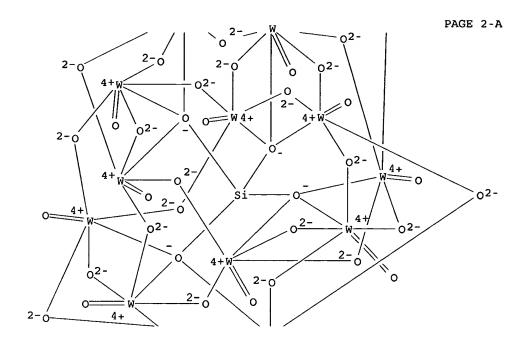


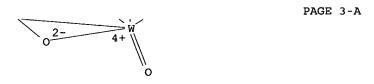




●3 H⁻¹

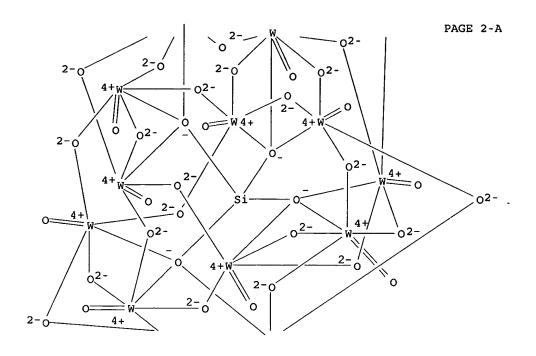


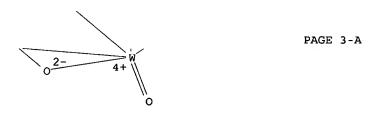




●4 H+

RN 12027-47-3 HCAPLUS
CN Tungstate(4-), [μ12-[orthosilicato(4-)κ0:κ0:κ0:κ0':κ0':κ0'
':κ0'':κ0'':κ0''':κ0''']]tetra
cosa-μ-oxododecaoxododeca-, tetrasodium (9CI) (CA INDEX NAME)



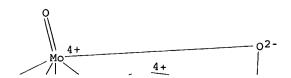


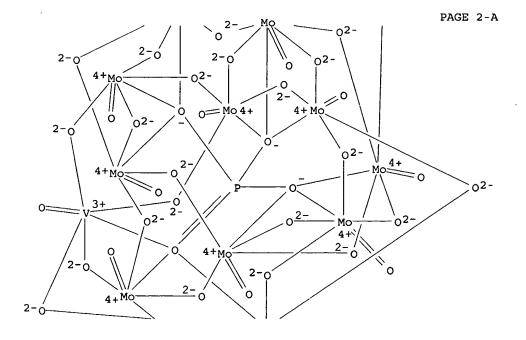
●4 Na+

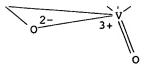
RN 12293-21-9 HCAPLUS

CN Vanadate(5-), (heptadeca-μ-oxodecaoxodecamolybdate)hepta-μ-oxodioxo[μ12-[phosphato(3-)-κΟ:κΟ:κΟ:κΟ
':κΟ':κΟ':κΟ'':κΟ'':κΟ'''
:κΟ''':κΟ''']]di-, pentahydrogen (9CI) (CA INDEX NAME)

PAGE 1-A

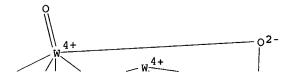


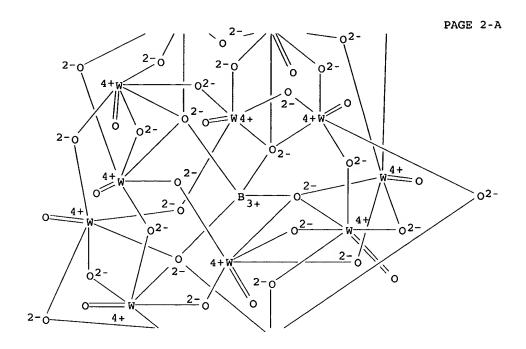


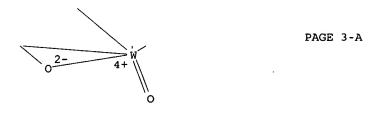


●5 H+

RN 12297-12-0 HCAPLUS
CN Tungstate(5-), tetracosa-μ-oxododecaoxo[μ12[tetrahydroxyborato(5-)-κΟ:κΟ:κΟ::.kapp
a.0':κΟ'':κΟ'':κΟ'':.kappa
.0''':κΟ''']]dodeca-, pentahydrogen (9CI) (CA INDEX NAME)



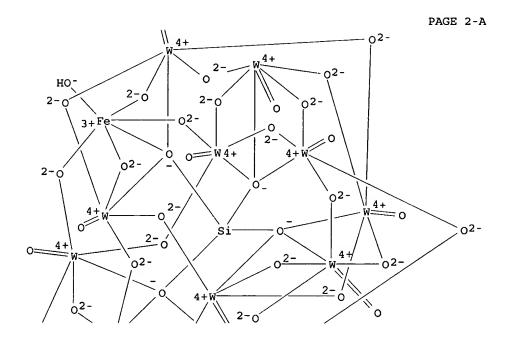


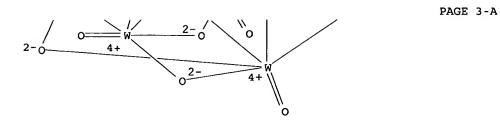


●5 H+

RN 37300-77-9 HCAPLUS CN Tungstate(6-), (hydroxyferrate) [μ 12-[orthosilicato(4-)-0:0:0:0':0':0':0'':0'':0''':0''']] tetracosa- μ -oxoundecaoxoundeca-, hexapotassium (9CI) (CA INDEX NAME)

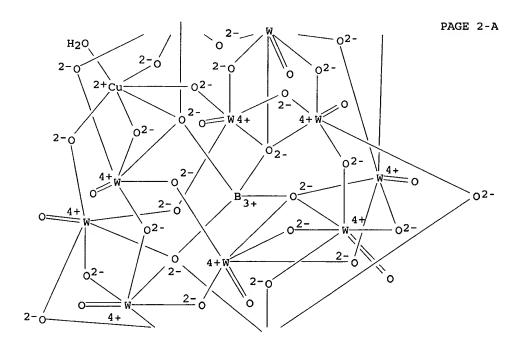


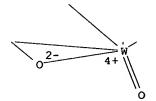




●6 K+

RN 37306-30-2 HCAPLUS
CN Tungstate(7-), (aquacuprate)tetracosa-μ-oxoundecaoxo[μ12[tetrahydroxyborato(5-)-O:0:0':0':0':0'':0'':0'':0''':0''']
]undeca-, tripotassium tetrasodium (9CI) (CA INDEX NAME)





●3 K+

●4 Na+

A61K; C01B

```
CC
     1-5 (Pharmacodynamics)
     Section cross-reference(s): 3
     12026-57-2 12027-38-2 12027-47-3
     12207-33-9 12293-21-9 12297-12-0
                                         12339-25-2
     12398-81-1 37300-77-9 37300-91-7 37300-94-0
     37300-95-1 37306-30-2 37306-31-3 37308-25-1 37308-26-2 56367-34-1 56449-73-1 56449-74-2
                                                           110294-54-7
     RL: BAC (Biological activity or effector, except adverse); BSU
     (Biological study, unclassified); BIOL (Biological study)
        (virucide)
L114 ANSWER 63 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN
                          1964:407695 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                          61:7695
ORIGINAL REFERENCE NO.:
                          61:1243e-h,1244a
                          Conditions for quantitative precipitation of
TITLE:
                          phosphorus as ammonium molybdophosphate
AUTHOR(S):
                          Archer, D. W.; Heslop, R. B.; Kirby, R.
CORPORATE SOURCE:
                          Univ. Manchester, UK
SOURCE:
                          Analytica Chimica Acta (1964), 30(5), 450-9
                          CODEN: ACACAM; ISSN: 0003-2670
DOCUMENT TYPE:
                          Journal
                          English
LANGUAGE:
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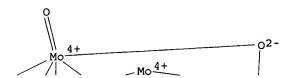
The conditions for quant. precipitating PO43- as ammonium molybdophosphate, (NH4)3(P(Mo3-O10)4), (I) were determined with resp. to temperature, time of standing, stirring, the MoO42-/PO43- ratio, the effect of HCl, H2SO4, HNO3, and HClO4, and addition of NH4NO3 before precipitation by measuring the β -activity of H3 32PO4 in the filtrates from I. The H332PO4 was prepared by mixing 2.5 ml. concentrated HCl, 1 ml. of 30% H2O2, 2.5 ml. of 0.0167 M KH2PO4, and 40 $\mu c.$ of 32P as H332PO4, warming on a hot plate for 15 min., and diluting to 100 ml. with H2O. The β -activity of filtrates (≥10,000 counts) was counted using a liquid counter and an automatic scaler. To determine the time and temperature for quant. precipitating PO43-, 4 ml. of nitromolybdate (II) (prepared according to Thistlethwaite, CA 42, 2206i) was added to 2 ml. of 0.0167 M KH2PO4, containing about 0.5 $\mu c.\ H3\ 32PO4\,,$ in a series of expts. at different temps. (30-90°). The solution was held at the selected temperature, stirred every 15 min., and allowed to stand for the same period of time at 25°. I was filtered, washed with acid NH4NO3, and finally with 1% HNO3. A β-count was made on the combined filtrate and washings and on the NH40H extract of I. The % PO43- precipitated was calculated from the corrected count rates. Mo, PO43-, and NH4+ were determined in I. After dissolving I in a min. volume of M NaOAc (pH 8.7), Mo6+ was determined as the 8-quinolinolate,

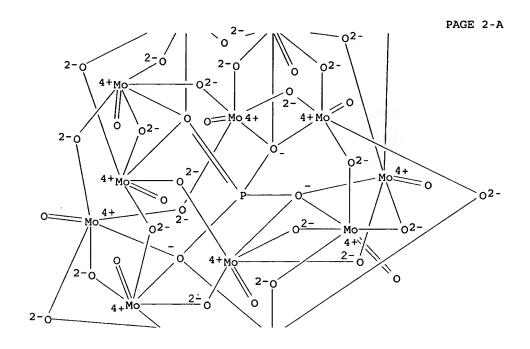
MoO2(C9H6ON)2 (III). The filtrate and washings from III were adjusted to pH 5.8 with M NaOH, and PO43- was determined as Pb3-(PO4)2. PO43- in I also was determined as MgNH4PO4.6H2O, after dissolving I in a min. volume of dilute NH4OH. NH4+ was determined by distillation of NH3 and titration. The factors of MoO42-/PO43, the effect of mineral acids, and addition of NH4NO3 were determined in a similar way. PO43- is precipitated quant. with 2-5.2 times the stoichiometric amount of II at 50 80°, allowing the mixture to stand for ≥30 min. at that temperature and 30 min. at 25° stirring at 15 min. intervals. The composition of the precipitate formed at 80° is between I and (NH4)2H(P(Mo3O10)4), as found by Thistlethwaite (loc. cit.). The precipitate formed at 90° has a Mo/P ratio > 12 and a NH4+ content less than that of I formed at 80°. HNO3 should be added; 3.8-6.5 M is optimum for precipitating I at 70 or 80°, but 10 M HNO3 allows quant. precipitation HClO4, 0.5-1.7 M, does not interfere, but HCl and H2SO4do. NH4NO3 has no effect. Peptization of the precipitate occurs if (NH4)2MoO4, HNO3, and PO43- are mixed sep.

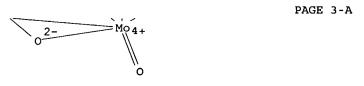
RN 12026-66-3 HCAPLUS

CN Molybdate(3-), tetracosa-μ-oxododecaoxo[μ12-[phosphato(3-)κ0:κ0:κ0:κ0':κ0':κ0'

':κΟ'':κΟ'':κΟ''':κΟ''']]dodec a-, triammonium (9CI) (CA INDEX NAME)





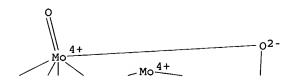


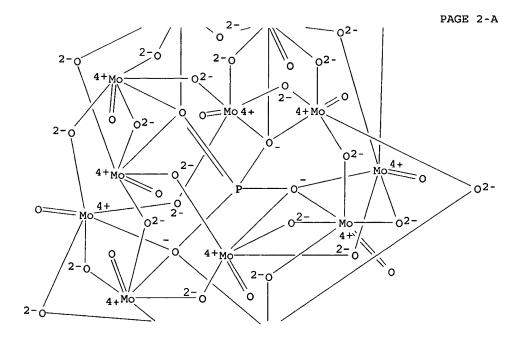
●3 NH4+

RN 12026-65-2 HCAPLUS

CN Molybdate(3-), tetracosa-µ-oxododecaoxo[µ12-[phosphato(3-)-0:0:0:0':0':0':0'':0'':0''':0''']]dodeca-, diammonium hydrogen (9CI) (CA INDEX NAME)

PAGE 1-A





2- Mo4+

PAGE 3-A

● H+

●2 NH4+

CC 2 (Analytical Chemistry)

IT 12026-66-3, Ammonium molybdophosphate, (NH4)3PMo12O40

12704-86-8, Ammonium molybdophosphate

(precipitation of)

IT 12026-65-2, Ammonium molybdophosphate, (NH4)2HPMo12O40 (preparation of)

L114 ANSWER 64 OF 64 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1929:20401 HCAPLUS

DOCUMENT NUMBER: 23:20401 ORIGINAL REFERENCE NO.: 23:2405d-h

TITLE: High-test cast iron

AUTHOR(S): Lemoine, R. P.

SOURCE: Am. Foundrymen's Assoc. (preprint) (1929), No.

29-12, 259-88

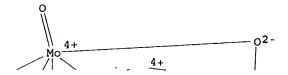
DOCUMENT TYPE: Journal LANGUAGE: Unavailable

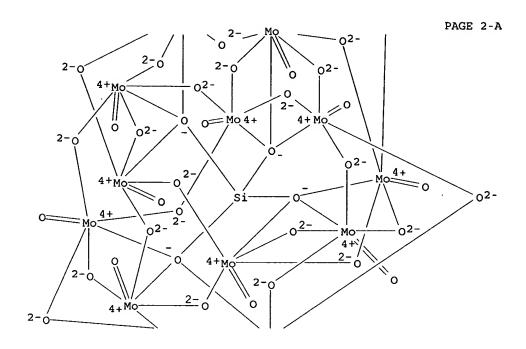
A graphical representation of the constitution of Cast Fe is discussed. The best mech. properties are attained when the Fe is composed only of graphite and pearlite, with the lowest total C.Free cementite is less detrimental than free ferrite. The cooling speed must be controlled to get equally good properties in different thicknesses of section. The Si content has somewhat the same influence as the cooling speed, but it also tends to diminish the C content of pearlite, so that under certain conditions increased Si giving more graphite does not involve the formation of ferrite, With low total C and high Si, the effect of cooling speed is diminished, giving uniformity of structure in all sections, and high strength. The French practice in making semi-steel shells during the war is reviewed. The Lanz and Thyssen-Emmel processes, based on, the pearlite-graphite structure, are not considered novel, or practical for varied work. The use of expensive alloys in cast Fe is not necessary for high mech. strength. Elec. melting is useful for the chemical improvement of poor raw materials, and for refining the graphite, possibly by allowing kish to float out at high temperature Steel scrap is the best base for elec. furnace charges, which can readily be brought by addns. to any desired composition The cost may be reduced by using a cupola for melting, and refining electrically. In this duplex practice, the cupola product should be as near the correct C content as possible, to save current consumption. To obtain a low-C cast Fe from the cupola, melting of steel scrap must be hastened by high airpressure with only 12 to 14% coke, so that the fused drops passing the tuy eres contain only 2% C. The use of a bottom receiver is an advantage to raise the temperature, and the absorption of too much C in the receiver is best prevented by

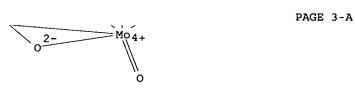
replacing some of the coke in it with brick fragments. Low-C Fe made in this way from rusty scrap may be wild and oxidized, but with ample Si it should be sufficiently fluid and quiet. Losses of Si through the cupola are lessened with an acid slag. High-Si steel scrap and high-Si pig Fe in the charge, are useful; pig Fe containing 2.7% C and 4% Si is practical and convenient, but addns. of ferric Si may be resorted to. The quality of such Cast Fe is between that of semi-steel and elec. cast Fe, and its cost is not excessive.

- IT 12027-12-2, Silicomolybdic acid, H4SiO4.12MoO3
 (effect on cast iron)
- RN 12027-12-2 HCAPLUS

NAME)







●4 H+

=>